



This Scanning Electron Micrograph (7000X) is the first 3-dimensional view of a cell in an ulcerated duodenum. The center is completely denuded, as the "best exhibit on original research or instruction on a medical subject" at the A.M.A. Clinical Convention, November 26-29, 1972, in Cincinnati, Ohio.

## The Tireless Man

### whose duodenal ulcer needs a rest

Up early, home late, often with a scratch pad filled with notes, figures, plans. A few hours' sleep and then another long day. This is often the routine of the tireless hard driver, one-man committee with enough overwork and stress to wear out several men. But his duodenal ulcer may warn him with sharp discomfort that he had better ease up, let some things go, and give himself—and his ulcer—a rest.

### The need to reduce G.I. hypermotility and hypersecretion

Overwork together with overanxiety are often principal factors in exacerbating a duodenal ulcer. To help reduce the increased gastric secretions and hypermotility, therapy may need to include treatment for associated undue anxiety—which is where dual-action Librax can be highly useful.

### The dual nature of Librax

Only Librax combines, in one capsule, the antianxiety action of Librium® (chlorodiazepoxide HCl) and the antisecretory action of Quarzan® (clidinium Br).

Before prescribing, please consult complete product information, a summary of which follows:

**Indications:** Symptomatic relief of hypersecretion, hypermotility, and anxiety and tension states associated with organic or functional gastrointestinal disorders; and as adjunctive therapy in the management of peptic ulcer, gastritis, duodenitis, irritable bowel syndrome, spastic colitis, and mild ulcerative colitis.

**Contraindications:** Patients with glaucoma; prostatic hypertrophy and benign bladder neck obstruction; known hypersensitivity to chlordiazepoxide hydrochloride and/or clidinium bromide.

**Warnings:** Caution patients about possible combined effects with alcohol and other CNS depressants. As with all CNS-requiring complete mental alertness (e.g., operating machinery, driving). Though physical and psychological dependence have rarely been reported on recommended doses, use caution in administering Librium (chlorodiazepoxide hydrochloride) to known addiction-prone individuals (those who convulse), following discontinuation of the drug and after withdrawal symptoms (including convulsions), following discontinuation of the drug and after discontinuation of the drug in pregnancy, lactation, or in women of child-

bearing age requires that its potential benefits be weighed against its possible hazards. As with all anticholinergic drugs, Precautions: In elderly and debilitated, limit dosage to small overdosage or confusion (not more than two capsules per day initially; increase gradually, as needed and tolerated).

With other psychotropics seems indicated, carefully considering drugs such as MAO inhibitors and phenothiazines. Observe usual precautions in patients with impaired renal or hepatic function. Paradoxical reactions (e.g., excitement, pallor, Employ usual precautions in treatment of anxiety states with evidence of impending depression; suicidal tendencies may be present and protective measures necessary. Variable effects on blood coagulation have been reported, rarely in patients receiving the drug and oral anticoagulants. Adverse Reactions: No side effects or manifestations not seen when chlorodiazepoxide hydrochloride is used alone, droperidol, and/or low residue diets.

### adjunctive Librax®

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

### For the anxiety-linked symptoms of duodenal ulcer

### Librax®

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br.

Each capsule contains 5 mg chlordiazepoxide HCl and 2.5 mg clidinium Br

## Ontario MDs' Medicare Records to Be Checked

Medical Tribune World Service

TORONTO—Ontario physicians who treat an unusually large number of patients under Medicare are going to have their records closely scrutinized by the College of Physicians and Surgeons of Ontario, the province's professional licensing authority.

The college announced here that any doctor who provides more than 300 units of service to patients in a week will be

subject to investigation. A unit, equal to an office visit, is worth \$6. A house call, at \$9, would be a unit and a half.

The Ontario Medical Association has agreed to set up a joint committee with the Government that would determine how much and how doctors should be paid.

Dr. Richard T. Potter, Ontario Minister of Health, said the Government has no intention of putting all doctors on salary. But he stressed that the ceiling set on a

physician's work load by the college demonstrates concern for the quality of medical care.

"As a family physician myself, I know that the doctor who is seeing more than 300 patients a week is getting damned little sleep and doing nothing else," Dr. Potter declared. He said that he "can't buy the claim" by one physician who billed the Medicare system for seeing 316 patients in one day.

## Breast-Feeding Has Dropped 49% in Sweden

Medical Tribune World Service

STOCKHOLM—From 1944 to 1970 the number of six-month-old Swedish babies being breast-fed dropped from 56 per cent to 7 per cent.

In an effort to discover why, Drs. Yngve Hofvander and Stig Sjölin, of the pediatric clinic at Uppsala's University Hospital, gave one-hour questionnaire interviews to 298 mothers of 302 children.

The investigators found that on discharge from the maternity ward, 72 per cent of the women were breast-feeding exclusively and 87 per cent partially. The frequency dropped rapidly after their return home.

Average duration of breast-feeding was nine weeks. By the end of the first month, 56 per cent had stopped completely or had introduced regular alternative feeding forms. By the end of the third month fewer than one-third were still breast-feeding and only one-sixth exclusively. At six months, 4 per cent were still breast-feeding for certain meals.

### 92% Planned to Breast-Feed

Some 92 per cent of the mothers said they had planned to breast-feed, most of them "as long as they could," at the time of delivery. Asked directly what they thought about breast-feeding, 68 per cent said they liked it "a lot" or "relatively much." Only 11 per cent admitted to not liking it.

The main reason for stopping reported by 66 per cent was that they "just ran dry." Only 10 per cent blamed "faulty breasts" or "deficiencies in their children."

"Most mothers are, or pretend to be, unaware of the real reasons for early discontinuance of breast feeding," Dr. Hofvander commented.

When asked why they thought other mothers discontinued breast-feeding early, the mothers provided answers that in many cases were at variance with those they applied to themselves.

## Japan Plans to Establish Continuing MD Education

Medical Tribune World Service

TOKYO—The Health and Welfare Ministry plans to establish a system of lifelong education for doctors. Ten professors are to be selected from the Japan Medical Association to draft the details of the program.

Under the proposed system, all physicians will be required to undergo regular qualification tests.

### NEWS INDEX

**Medicine:** pgs. 1, 2, 3, 7, 9, 29, 31, 32, 33, 35, 38, 39

FDA is charged with blocking the introduction of valuable new drugs for migraine and headache ..... 9

Medical ethics teaching is said to be having a phenomenal growth in the curriculums of medical schools ..... 31

Hemorrhage duodenitis should be considered in the differential diagnosis of upper gastrointestinal bleeding ..... 33

Sensitivity to the cockroach may be a factor in causing allergic illnesses in cockroach-infested homes ..... 35

Physical and mental fatigue, for example, was given as a reason for cessation in others by 17 per cent, against 8 per cent for cessation by themselves. Also, 27 per cent thought others considered breast-feeding "messy," while only 5 per cent thought so themselves. Similarly, 38 per cent thought others considered breast-feeding to be uncomfortable and an encroachment on personal freedom, whereas only 4 per cent felt this themselves.

## Mercury Pollution Warnings Send Sale of Fish Tumbling

Medical Tribune World Service

KYUSHU, JAPAN—Warnings by the Government of the dangers of mercury pollution have sent fish sales tumbling in markets throughout Japan and created storage problems for fishing fleets and distributors.

Tentative tolerance levels for mercury in fish and shellfish have been set at the world's lowest levels, on the basis of a report by a group of experts set up after the third confirmed outbreak of Minamata disease in Kyushu in May.

The weekly intake for an adult has been fixed at 0.17 mg. of methyl mercury. The average concentration of all mercury must be below 0.4 ppm, and 0.3 ppm in the case of methyl mercury, compared with 0.5 ppm of all mercury set by the United States and Canada and 1 ppm by Sweden and Finland. The experts warned that pregnant women, in particular, should strictly observe the levels indicated because of the susceptibility of the fetus.

The Government has also issued a typical menu in its efforts to get citizens to hold their fish and shellfish intake to 567 gm. per week. One example for a week's consumption listed four small horse mackerel, half of a medium flatfish, and one medium squid. Another example was three young punctatus, one mackerel pike, one prawn, and 20 slices of raw tuna.

Health experts warned further that the

Only 0.5 per cent of the mothers offered personal appearance as a reason for having terminated breast-feeding. But 12 per cent believed this to have been the reason among other women.

"It is probable," Dr. Hofvander said, "that the mothers' conception of why others terminate breast-feeding to a certain, perhaps a great, extent is an unconscious projection of their own experiences, and thus best reflects their own feelings."

## Highest Cancer Mortality Found in 35-64-Year-Olds

Medical Tribune World Service

567-Gm. weekly limit refers to dried fish, fish meal, canned fish, and ham and sausage made from fish.

But, after the initial scare, the Government's Environmental Agency came out with a revised list of fish quantities—about double the first list—that could be safely consumed in a week.

## Airplane Exhaust Fumes Are Called Major Cause Of Pollution in Japan

Medical Tribune World Service

YOKOHAMA, JAPAN—Exhaust fumes from aircraft, not automobiles, are the major cause of photochemical smog in Japan, according to Prof. Tetsuzo Kungawa, of Yokohama National University.

The large quantities of exhaust fumes discharged by jet planes flying through the inversion layer become oxidized by strong rays of the sun, according to Professor Kitagawa's theory. His studies found that the exhaust fumes combine with vapor and they become three times heavier than the air. This mass then descends to just above ground level, causing localized photochemical smog.

He found that the smog usually occurs near airfields, especially in cities located between 35 and 45 degrees north latitude, which are exposed to the sun's rays at certain angles. This explains, he said, why the smog frequently occurs in areas where vehicle traffic is relatively light.

A DC8 aircraft, he said, consumes 25,800 L. of fuel when climbing, about 4,000 times the average amount consumed by a car.

CLINICAL NEWS NOTE: "Fiberoptic colonoscopy constitutes a real advance in the diagnostic and therapeutic armamentarium of the medical profession." (Dr. William L. Wolff; see page 33.)

**Ob/Gyn:** pgs. 1, 3, 4, 7, 8, 31, 32  
Teen-age mothers and their infants have a higher death rate, reflecting social consequences of teen pregnancy ..... 32

**Pediatrics:** pgs. 1, 6, 7, 35, 39, 40  
Growth-accelerating effect of chorionic gonadotropin is detectable within three weeks ..... 6

**Surgery:** pgs. 5, 6, 33, 43  
Fiberoptic colonoscopy requires adequate training programs for physicians to gain experience in performing the procedure under supervision ..... 33

**Psychiatry:** pgs. 1, 5, 9, 40  
Incestuous unions between fathers and daughters in Sweden are reportedly more common today than in the past ..... 5

## Herbs for Healing Sought



In Red China, where herbs are still often used for medicinal purposes, finding and picking them can be quite a job. Here climbers go up the side of Mount Huangshan in Anhwei Province to gather the valuable herbs. The area is known for its beautiful landscapes.

## Highest Cancer Mortality Found in 35-64-Year-Olds

Medical Tribune World Service

TOKYO—The heaviest cancer mortality in Japan in 1971 was in the 35-to-64-year age group. Cancer was responsible for 21.1 per cent of deaths in the age bracket of 35 to 39 years and 24.8 per cent in those from 40 to 44 years of age.

Stomach cancer still claimed the most victims, but the number of deaths is decreasing while lung cancer deaths are climbing.

## 1973 Smallpox Incidence Doubles in Comparison With First Half of 1972

Medical Tribune World Service

GENEVA, SWITZERLAND—The reported incidence of smallpox almost doubled during the first half of this year over that during the same period in 1972, the World Health Organization reported.

India and Bangladesh, where the disease is endemic, account for almost 88 per cent

of the 77,984 cases reported this year. Both countries have been reporting the highest incidence since the global program for eradication of smallpox began seven years ago.

Cases also occurred in four other countries of Asia this year. In Pakistan smallpox is also endemic, but all cases in Afghanistan (14), Japan (1), and Nepal (104) were the result of importation from India, Pakistan, and Bangladesh.

In Africa, WHO reported, progress con-

## Advice for the Chronically Ill



## MDs Told to Set Own Standards Or Lose Control

Medical Tribune Report

NEW YORK—The Federal Government's top physician reminded the American Medical Association here that if the medical profession does not set the standards for medical care, the job will be done by "some bureaucratic agency in Washington."

Dr. Charles C. Edwards, assistant secretary for health in the Department of Health, Education, and Welfare, told the A.M.A. House of Delegates that "the public is not going to accept a continuation of things as they are." He emphasized that "if physicians continue to insist on the freedom to exercise their own professional judgment, then they must accept the responsibility to assure the quality of the care they provide."

He said the A.M.A. was "making the right choice" by taking an active role in development of the Peer Standards Review Organization concept that came into law last fall with H.R. 1. He pointed out there were other activities that the A.M.A. might into.

For instance, said Dr. Edwards, the recent passage of legislation extending many Federal health programs for another year provides "a challenge and an opportunity to take another look at . . . Regional Medical Programs, Hill-Burton, Community Mental Health Centers, and the rest."

Dr. Edwards conceded that the Government had been "too long without a clear, articulate national health strategy . . . because the Federal health enterprise has not been organized to develop one." However, "we are now in the process of putting our own house in order," he said, "for the coordination of our efforts and for productive communication with those outside Government."

## Pulmonary Edema May Still Follow Drowning Rescue

Medical Tribune Report

NEW YORK—The risk of death from drowning does not end once the victim has been revived, a leading investigator warned here, adding that all near-drowning victims who require artificial respiration should be hospitalized for 24 hours immediately after the accident.

Lung damage can occur even if water is not breathed into the lungs, said Dr. Martin J. Nemiroff, of University Hospital, Ann Arbor, Mich. The brief period of suffocation and lack of oxygen during submersion can cause pulmonary edema, he warned the American Lung Association.

He stressed that near-drowning and related deaths are not recorded by law and thus, he suggested, their incidence may be more common than realized.

### Eight Cases Described

Dr. Nemiroff described eight cases with one death in one geographic area. Six of the eight walked away from the scene of the accident, only to be hospitalized after rapidly progressive shortness of breath two and a half to 12 hours later. Another was hospitalized immediately, and one died a half hour after near-drowning.

The amount of fluid in the lungs of the seven survivors varied, but increased in all of them during the first 24 hours after the near-drowning. The edema led to hypoxia, hypercapnia, and metabolic acidosis. The near-drowning also caused damage to the lung capillaries.

Continuous, positive-pressure breathing was used to prevent both lung collapse and the seeping of fluid into the lungs. Three patients whose x-rays showed the greatest amount of fluid in the lungs also required endotracheal intubation.

## Federal Safeguards Urged Against Sterilization Abuses

Medical Tribune Report

NEW YORK—The Planned Parenthood Federation of America has called on the Government to set up a working conference of experts to help draft Federal safeguards against "potential abuse" of contraceptive sterilization in persons who are "uneducated, young, emotionally immature, or mentally retarded."

Condemning the sterilization of several young girls by the Montgomery County Community Action Agency in Alabama, Dr. Allan F. Guttmacher, federation president, said: "In our view it is imperative that the Department of Health, Education, and Welfare issue regulations or guidelines to govern the provision of sterilization in Federal programs."

### Views Given in Letter

His views were made known in a letter to Dr. Louis Hellman, deputy assistant secretary for population affairs.

Dr. Guttmacher reaffirmed the federation's support of voluntary sterilization "for anyone who is fully informed of the nature of the procedure and who is mature

"Twenty-Fifth Wedding Anniversary dates are on the calendar in May and June for several members and warmest wishes and congratulations go to . . . —Dr. of the Lehigh County (Pa.) Medical Society. Those the odds, or what? (Regular beats Immateria Medica, page 43.)



## Bobo's back at the big top

For full details please read the prescribing information. It's summarized on the back of this page.

**Butazolidin® alka**  
Each capsule contains:  
100 mg. phenylbutazone USP  
100 mg. dried aluminum hydroxide gel USP  
100 mg. magnesium trisilicate USP

If it doesn't work in a week, forget it.

Geigy

MEDICAL TRIBUNE is published each Wednesday except on Jan. 31, May 30, Aug 29, and Oct. 31, by Medical Tribune Inc., 880 Madison Ave., New York, N.Y. 10021. Contracted circulation postage paid at Farmingdale, N.Y. 11735. Subscription \$12.50, Students, \$7.50.

## CLINIC OF THE MONTH

What's new and important in perinatal medicine?



## The Consultant

DR. ROBERT J. LUBY  
Department of Obstetrics and Gynecology,  
Creighton University, St. Joseph Hospital,  
Omaha, Neb.

... a significant reduction in perinatal mortality remains to be achieved in this country....

THE NEW AND IMPORTANT DEVELOPMENTS in perinatology may be considered in three categories: first, laboratory studies; second, equipment; and third, the impact of inhouse newborn specialists on observed newborn death rates.

First, the ability to study amniotic fluid phospholipid levels and to predict, with reasonable accuracy, those infants with pulmonary maturity has been most helpful.

Second, the ability to study amniotic fluid phospholipid levels and to predict, with reasonable accuracy, those infants with pulmonary maturity has been most helpful. Since Gluck's original report, we have used his exact technique in our laboratory and have found his results to be reproducible. To date, we have seen no infants die of respiratory distress with a mature pattern. We have encountered severe respiratory distress with an inter-

mediate pattern and with immature patterns, and infants with immature patterns who did not develop respiratory distress.

In the area of equipment, the availability of accurate, relatively low-cost

monitoring equipment for labor and delivery units has improved significantly the quality of observation of the infant during labor. The expenditure of \$7,000 to \$8,000 per unit, while significant, pales when one considers the cost of new diagnostic or therapeutic radiology equipment. This monitoring, plus the demonstration and availability of positive pressure treatment for the neonate, is extremely promising.

Dr. Graeven and others have shown repeatedly, insofar as the third category is concerned, that infant death rates are reduced from 15 to 17 per thousand live births to under 10 per thousand live births with the presence of a newborn specialist and his supporting skills in the delivery unit. It is obvious that prompt attention to required ventilatory and chemical resuscitation is rewarded in terms of useful life preservation.

When should amniotic fluid be analyzed to determine fetal maturity and how is this done?

Amniotic fluid analysis for purposes of fetal maturity determination should be performed whenever this information contributes to the management of the patient, e.g., pregnancies to be terminated by elective C-section when the estimated date of



## Butazolidin® alka

Each capsule contains:  
100 mg. phenylbutazone USP  
100 mg. dried aluminum hydroxide gel USP  
150 mg. magnesium trisilicate USP

A tough act  
to follow.

**Important Note:** This drug is not a simple analgesic. Do not administer casually. Carefully evaluate the patient before starting treatment and keep them under close observation. Obtain a detailed history and complete physical and laboratory examination (complete hemogram, urinalysis, etc.) before prescribing and at frequent intervals thereafter. Carefully select patients, avoiding those responsive to relief measures, contraindicated patients and those who should be observed frequently. Warn patients not to exceed recommended dosage. Short-term relief of severe symptoms with the smallest possible dosage is the goal of therapy. Dosage should be taken with meals or a full glass of fluid. Substituted alka capsules for tablets if possible. Do not exceed 400 mg. daily. Do not take any sign of: fever, sore throat, oral lesions (dry mouth or blood dyscrasias); diarrhea, epigastric pain, symptoms of anemia, black or tarry stools or other evidence of intestinal ulceration or hemorrhage; skin reactions, including hives, rash or edema. A one-week trial period is suggested. Discontinue in the absence of a favorable response. Restrict treatment periods to one week in patients over 60.

**Indications:** Acute gouty arthritis, rheumatoid arthritis, rheumatoid spondylitis.

**Contraindications:** Children 14 years of age; severe hypertension; history or symptoms of G.I. inflammation; history of peptic ulcer, severe, recurrent or peristaltic dyspepsia, diarrhea or presence of drug after or blood dyscrasias; renal, hepatic or cardiac dysfunction; hypertension; thyroid disease; systemic edema; anuria and salivary gland enlargement due to tumor; polymyalgia rheumatica and temporal arteritis; patients receiving other potent chemicotherapeutic agents, or long-term anticoagulant therapy.

**Precautions:** This is a potent drug. Its misuse can lead to serious results. Review detailed information before beginning.

**Adverse Reactions:** This is a potent drug. Its

therapy. Ulcerative colitis, acute and recurrent gastritis and duodenitis, ulcer with perforation and hemorrhage, ulceration and perforation of large bowel, occult G.I. bleeding with anemia, headache, epigastric pain, hematemesis, dyspepsia, diarrhea, vomiting and diarrhea, abdominal distension, dyspepsia, epigastric pain, hematochezia, anemia due to blood loss including occult G.I. bleeding, thrombocytopenia, pancytopenia, leukemia, leukopenia, bone marrow depression, sodium and chloride retention, water and sodium retention, plasma dilution, respiratory distress, renal colic, edema, fetal and neonatal hepatitis (which may or may not be prominent), petechiae, purpura without thrombocytopenia, toxic pruritis, erythema nodosum, erythema multiforme, Stevens-Johnson syndrome, Lyell's syndrome (toxic necrolytic epidermolytic), exfoliative dermatitis, serum sickness, hypotension, hypotension (polyarteritis), anaphylactic shock, urticaria, peripheral edema, rash (allergic reactions requiring parenteral and permanent withdrawal of the drug), proteinuria, hematuria, oliguria, anuria, renal failure with azotemia, glomerulonephritis, acute tubular necrosis, nephritic syndrome, bilateral renal cortical necrosis, renal stones, ureteral obstruction with uric acid crystals due to overexcretion of drug, impaired renal function, edema, hypertension, hypertension, periorbital, diffuse, periorbital myocardiitis with muscle necrosis, periorbital granulomatosis, aggravation of temporal arteritis in patients with polymyalgia rheumatica, optic neuritis, blurred vision, retinal hemorrhage, toxic amblyopia, retinal detachment, hearing loss, hyperglycemia, hyperglycemia, toxic goiter, association of hypertension and hypotension (causal relationship not established), agitation, confusion, states, lethargy, CNS reactions associated with overdose. Including convulsions, euphoria, psychosis, depression, headaches, hallucinations, giddiness, vertigo, com. hyperventilation, insomnia, tachycardia, salivation, salivary gland enlargement. (B10-146-070-H10/71)

For complete details, including dosage, please see full prescribing information.

**GEIGY Pharmaceuticals**  
Division of CIBA-GEIGY Corporation  
Ardsley, New York 10502

© 1973 CIBA-GEIGY Corporation

GEIGY



## A.M.A. Opponents Of Abortion Fail To Reverse Policy

Medical Tribune Report

**NEW YORK**—The forces against abortion surfaced again here at the annual convention of the American Medical Association in an effort to offset the organization's 1970 policy relaxation that mainly says "abortion is a medical procedure."

Four California resolutions tackled the issue in ways that ranged from decrying the lack of children available for adoption to taking "a positive view of motherhood." A Louisiana resolution wanted state legislatures to adopt an amendment to the U.S. Constitution that would make abortion a matter of states' rights.

A committee of the A.M.A. House of Delegates that conducted hearings on the California measures said that it received more than a hundred telegrams and other messages on the issue, plus "an official statement from the Committee of Doctors and Nurses Against Abortion."

However, said the house committee, the existing A.M.A. policy provides that medical personnel shall not be compelled to perform abortions in violation of their medical judgment or moral principles. The committee recommended reaffirmation of the 1970 policy.

And the delegates reaffirmed it, in spite of the contention by Dr. Joseph P. Donnelly, delegate from Newark, N.J., that "our present policy favors unlimited abortion."

Immediately afterward the house also adopted a statement "affirming the traditional favorable attitude of the medical profession toward pregnancy and motherhood." This entails the establishment of "counseling programs that will offer constructive help to expectant mothers in accepting and coping with the stresses of pregnancy" and provides "incentives such as approval, appreciation, encouragement, and emotional support for a decision to continue pregnancy to term."

### Amendment Called Premature

On the resolution for a U.S. Constitutional amendment, another house committee decided it was "premature" and suggested instead that the A.M.A. "monitor and study" the effects of the recent Supreme Court ruling that wiped out all state prohibitions against abortion. The delegates agreed to that approach.

But they were not finished with normalize issues, and the two following items proved less easy to resolve. Both were introduced by the now Interns and Residents Section of the A.M.A.

One asked that the organization work to end legal and employment discrimination against homosexuals and legal restrictions on sexual behavior between consenting adults. The house sent the idea to the A.M.A. Council on Mental Health.

The other resolution sought A.M.A. endorsement of a program that would not only "teach moral and social responsibility" to youngsters but also push for state laws to "allow condom contraceptives to be displayed and sold openly above the counter without age restrictions."

A Florida delegate took the floor to ask sarcastically whether the house also should approve "IUDs at Girl Scout meetings, diaphragms in cereal boxes, and 'the pill' in bubble gum dispensers."

The delegate of the interns and residents, Dr. Eugene S. Osgood, said he had to "regret the moral focus . . . [when] the issue is venereal disease, not pregnancy or morals."

Dr. Donnelly saw the item as part of "a great moral breakdown" and warned that "it won't solve our problems anymore than other permissive actions have." He scorned the discussion itself, saying, "Goddamn it, once upon a time this organization had a moral conscience."

The delegates tabled the matter until it comes up again. The next opportunity will be the November convention in Anaheim, Calif.

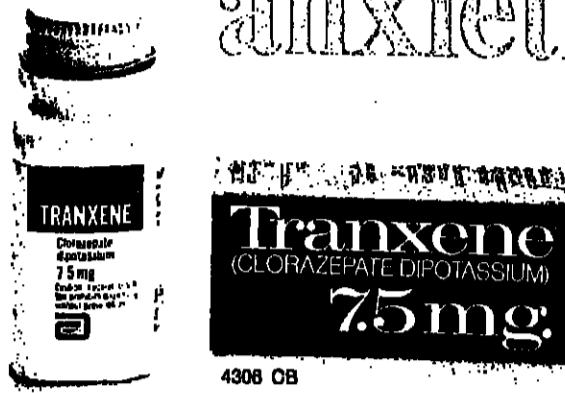
Wednesday, August 8, 1973

Wednesday, August 8, 1973

MEDICAL TRIBUNE

## No panacea. No placebo. No antidote for the pressures of everyday living.

## But a drug help relieve crippling anxieties



Tranxene has just one purpose: to offer effective control of symptoms for the patient with clinically manifested anxiety.

—*the patient whose anxieties are excessive and "inappropriate" to the circumstances at hand*

—*the patient with persistent (and often inexplicable) feelings of dread*

—*the patient who reacts unreasonably to reasonable stresses, to the point of incapacitation*

—*the patient with a sense of impending death or catastrophe (often seen as a complication of organic illness, such as cardiac disease)*

—*the patient with the physical symptoms of acute anxiety: sweating, insomnia, extreme nervousness, palpitations*

### Effectiveness shown in double-blind studies

The clinical investigation of Tranxene took place over four years; treatment periods ranged from

three week to six months.

A total of 50 efficacy studies were conducted, under controlled, double-blind conditions. The overall results showed Tranxene to be highly effective in relieving the symptoms of anxiety.

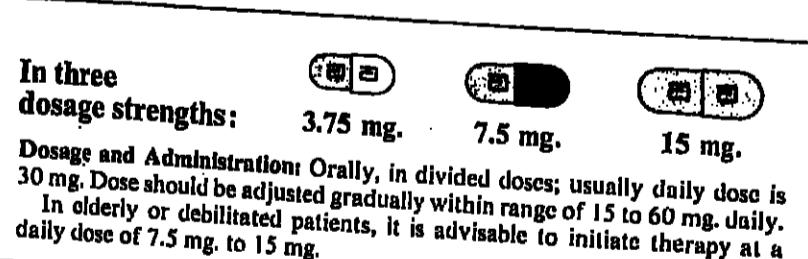
### Well tolerated by patients

Tranxene has an excellent record of patient acceptance. In the clinical studies, serious adverse reactions were not seen at the recommended doses. The side effects most commonly reported were drowsiness, light-headedness and gastrointestinal complaints.

### Minimal cardiovascular effects

In the clinical studies, the only effect seen on blood pressure was the lowering of slightly elevated systolic blood pressure in some patients. There were no reports of bradycardia and, in the two studies where electrocardiographic effects were studied, no evidence of drug-induced alterations in ECGs.

Where anxiety symptoms must be controlled, Tranxene can be a valuable—and prudent—aid in management.



In three dosage strengths: 3.75 mg. 7.5 mg. 15 mg.

Dosage and Administration: Orally, in divided doses; usually daily dose is 30 mg. Dose should be adjusted gradually within range of 15 to 60 mg. Daily, daily dose of 7.5 mg. to 15 mg.

DESCRIPTION: Chemically, TRANXENE (clorazepate dipotassium) is a benzodiazepine. The empirical formula is  $C_{14}H_{11}ClN_2O_2$ ; the molecular weight is 408.92. The compound occurs as a fine, light yellow, practically odorless powder. It is insoluble in the common organic solvents, but very soluble in water. Aqueous solutions are unstable, clear, light yellow, and alkaline.

ACTIONS: Pharmacologically, TRANXENE (clorazepate dipotassium) has the characteristics of the benzodiazepines. It has depressant effects on the central nervous system. The primary metabolite, nordiazepam, reaches peak level in the blood stream at approximately 1 hour. The plasma half-life is about 1 day. The drug is metabolized in the liver and excreted primarily in the urine. (See ANIMAL AND CLINICAL PHARMACOLOGY)

INDICATIONS: TRANXENE is indicated for the symptomatic relief of anxiety associated with anxiety neuroses, in other psychoneuroses in which anxiety symptoms are prominent features, and as an adjunct in disease states in which anxiety is manifested.

CONTRAINDICATIONS: TRANXENE (clorazepate dipotassium) is contraindicated in patients with a

Caution should be observed in patients who are considered to have a psychological potential for drug dependence.

Evidence of drug dependence has been observed in dogs and rabbits which was characterized byulsive seizures when the drug was abruptly withdrawn or the dose was reduced; the syndrome in dogs could be abolished by administration of clorazepate.

Usage in Pregnancy: Reproduction studies have been performed in rats and rabbits and there was no evidence of harm to the animal fetus. The relevance to the human is not known. Since there is no experience in pregnant women who have received this drug, safety in pregnancy has not been established.

It is assumed that TRANXENE or its metabolites is excreted in human milk. Therefore, this drug should not be given to nursing mothers.

PRECAUTIONS: In those patients in which a degree of depression accompanies the anxiety, suicidal tendencies may be present and protective measures may be required. The least amount of drug that is feasible should be available to the patient.

Patients on TRANXENE for prolonged periods should have blood counts and liver function tests periodically.

The usual precautions in treating patients with impaired renal/hepatic function should also be observed. In elderly or debilitated patients, the initial dose should be small, and increments should be made gradually, in accordance with the response of the patient, to preclude ataxia or excessive sedation.

ADVERSE REACTIONS: This side effect most frequently reported was drowsiness. Less commonly reported (in descending order of occurrence) were abrupt discontinuance of clorazepate, symptoms of nervousness, insomnia, irritability, dizziness, muscle aches and memory impairment, have followed abrupt withdrawal after long-term use of high dosage.

MANAGEMENT OF OVERDOSAGE: As in the management of overdose with any drug, it should be

simi skin rashes, fatigue, ataxia, genito-urinary complaints, irritability, diplopia, depression and slurred speech.

There have been reports of abnormal liver and kidney function tests and decrease in hematocrit.

Decrease in systolic blood pressure has been observed.

USAGE AND ADMINISTRATION: TRANXENE is administered orally in divided doses. The usual daily dose is 30 mg. The dose should be adjusted gradually, within the range of 15 to 60 mg, daily. In accordance with the response of the patient, doselessness may occur at the initiation of treatment patients. It is advisable to initiate treatment in elderly or debilitated patients.

DRUG INTERACTIONS: If TRANXENE (clorazepate dipotassium) is to be combined with other drugs acting on the central nervous system, careful consideration should be given to the pharmacology of the agents to be employed. Animal experiments indicate that TRANXENE prolongs the sleeping time after hexobarbital or other ethyl alcohol increases the inhibitory effect of clorazepate, but does not exhibit monoamine oxidase inhibition. Clinical studies have shown increased sedation with concurrent hypnotic medications.

The actions of the benzodiazepines may be potentiated by barbiturates, narcotics, phenothiazines, monoamine oxidase inhibitors, other anti-depressants, with somatic disease states, careful attention must be paid to possible drug interaction with concomitant therapy.

HOW SUPPLIED: TRANXENE (clorazepate dipotassium) is supplied as capsules in three dosage strengths: 3.75 mg. capsules (gray with white cap) in bottles of 100 (NDC 074-3417-19) and 500 (NDC 074-3417-53). 7.5 mg. capsules (gray with maroon cap) in bottles of 100 (NDC 074-3418-13) and 500 (NDC 074-3418-53). 15 mg. capsules (all gray) in bottles of 100 (NDC 074-3419-13) and 500 (NDC 074-3419-53).

Examination of all organs revealed no alterations attributable to TRANXENE. There was no damage to liver function or structure.

Reproduction Studies: Standard studies of fertility, teratology and reproduction were conducted on rats and rabbits. Oral doses in rats up to 150 mg./kg. and in the females up to 15 mg./kg. produced no abnormalities in the fetuses and no impairment to fertility and reproductive capacity of adult animals attributable to TRANXENE (clorazepate dipotassium). As expected, the sedative effect of high doses interfered with care of the young by their mothers (see Use in Pregnancy).

Clinical Pharmacology: Studies in healthy men have shown that TRANXENE has depressant effects on the central nervous system. Prolonged administration of high doses (120 mg. daily as a single oral dose) was without toxic effects, and abrupt cessation of drug was not followed by serious signs or symptoms.

Absorption—Excretion: After oral administration of TRANXENE (clorazepate dipotassium), there is essentially no circulatory parent drug. Nordiazepam, its primary metabolite, quickly appears in the blood stream with peak levels at about 1 hour. The plasma half-life is approximately 1 day. In volunteers given 15 mg. (50  $\mu$ C) of  $^{14}$ C-Tranxene, about 80% was recovered in the urine and feces within 10 days. Excretion was primarily in the urine with about 15% excreted per day on day 10.

Twenty-four dogs were given TRANXENE orally in a 22-month toxicity study; doses up to 75 mg./kg. were given. Drug-related changes occurred in the liver: weight was increased and cholestasis with minimal hepatocellular damage was found, but lobular architecture remained well preserved.

Eighteen rhesus monkeys were given oral doses of 100 mg./kg. daily for 52 weeks. All treated animals remained similar to control animals. Although the neutrophil count remained within normal limits, it tended to fall in the female animals on the highest doses.

## Headache Expert Charges FDA Blocks Valuable New Drugs

Medical Tribune Report

**NEW YORK**—The president of the American Association for the Study of Headache charged here that the Food and Drug Administration has blocked the introduction of valuable new drugs for migraine and headache.

Asserting that the effort to "ensure safety for all is to deny therapy to any," Dr. Seymour Diamond declared that only five new prescription drugs were introduced in the United States in 1970, and "none of the 75 pharmaceuticals which were introduced in England between 1966 and 1972 have been approved for use in the United States."

He noted that pizotyline and clonidine have been found effective for use in some patients with migraine and headache, but he stressed that the drugs are not available to American physicians.

"Unwarranted" obligations placed upon the FDA by Congress, Dr. Diamond said, "have caused years of delay in the introduction of such excellent drugs as metronidazole, levodopa, lithium and another half-dozen fine products."

The FDA's reluctance to license new drugs has often forced physicians to use existing compounds in spite of FDA rules limiting their use, he asserted.

"All of us have used drugs such as propantheline, or the MAO inhibitor, or large doses of ergotamine tartrate in the prophylaxis of migraine. We have used these drugs in spite of the package circular or description of the drug which limits its use to other purposes or limits the amount of drug used to an insufficient amount to help the patient."

He added that the exclusion of certain drug uses "intimidates the headache practitioner and leaves him liable to malpractice suits."

## Self-Help Units Win Growing Acceptance By Psychiatrists

Medical Tribune Report

**CHICAGO**—A survey of persons participating in group meetings of Recovery, Inc., a national self-help organization of nervous or former mental patients, has disclosed that more than one-third are attending the self-help sessions on the advice of professional counselors.

The influence exerted by referrals from psychiatrists, family physicians, social workers, religious advisers, and other professionals was reported by Dr. Hanus J. Grosz, Professor of Psychiatry, Indiana University School of Medicine, Indianapolis, following a survey of 6,463 members among 500 Recovery groups.

Dr. Grosz commented that acceptance of psychologic self-help by psychiatrists reflects a growing trend toward endorsement of groups run by and for persons seeking to overcome particular problems or to gain strength from mutual support so that they do not relapse.

Thirty-seven per cent of those surveyed were found to have been referred to Recovery, Inc., by a professional adviser. A psychiatrist's advice accounted for 20 per cent of these referrals.

In addition to the professional referrals, Dr. Grosz found that 23 per cent were referred by other Recovery members, 22 per cent by friends, and 15 per cent by relatives.

### Obesity, Drinking Affect Diving

**TOKYO**—Overweight skin divers and those who drink heavily are more likely to suffer from air embolisms, medical researchers at the Kitakyushu Workers' Accident Hospital reported. Dr. Kou Hayashi said a study of 300 skin divers showed that 90 per cent of the overweight divers and 95 per cent of the frequent drinkers have a history of the bends.

# NEW Intal® capsules 20 mg. cromolyn sodium, Fisons the one you've been hearing about.

Now available for your prescription in  
the United States.

Intal Caps  
#60  
with Spinhaler  
Sig: 1 capsule  
by inhalation  
q.i.d.

A Fisons' Representative will  
be calling on you soon to tell you  
more about this unique new discovery.  
Also, watch your journals and mail  
for more details.

Note: Spinhaler  
need only be specified  
on initial Rx.

**FISONS CORPORATION**

Bedford, Massachusetts 01730

Wednesday, August 8, 1973

MEDICAL TRIBUNE

11

## Equal Abortion Rights Are Urged for Minors

Medical Tribune Report

**SAN FRANCISCO**—The American College of Legal Medicine has urged equal rights for minors in an abortion resolution adopted at its 13th International Conference on Legal Medicine here.

The resolution asserts that the college's previous position on abortion is to be applied equally to minors and adults.

"In addition, where state law demands parental consent," the resolution says, "it is possible for the parents to effectively obviate the wishes of the pregnant minor. The college recommends that a procedure be established whereby a minor can seek the assistance and consent of the courts within a short period of time following the parents' adverse decision."

**Court Decisions Cited**

In another resolution the college cited "recent court decisions awarding damages to relatives of a decedent where an autopsy was performed as authorized by one relative but unknown to other relatives." It recommended, therefore, that "legislation be enacted to authorize any single relative, guardian, or legal representative of a decedent to consent to a postmortem ex-

amination and autopsy on a decedent's body for the purpose of determining the cause of death, for the advancement of medical or dental education and research, and for the general advancement of medical or dental science, provided that no person in a higher class exists or all persons in a higher class are not reasonably available at the time of death."

The college proposed that the priority of classes be as follows: (1) the spouse; (2) an adult son or daughter; (3) either parent; (4) an adult brother or sister; (5) a guardian of the person of the decedent at the time of death; or (6) any other person who has been authorized or is under

legal obligation to dispose of the body. The college also supported a resolution presented by Dr. R. L. Sadoff of Jenkins, Pa., appealing to all elected and appointed officials to uphold the law to protect the privacy of all patients, and one offered by Dr. Herman Wing of Chicago, which stated:

"The American College of Legal Medicine condemns the infringement and encroachment on the confidentiality of physicians' records even in their private offices, resulting from government proposals and third-party intrusions. This is a violation of proper medical-legal principles of the doctor-patient relationship."

## Review of Neurosurgery Training Is Planned

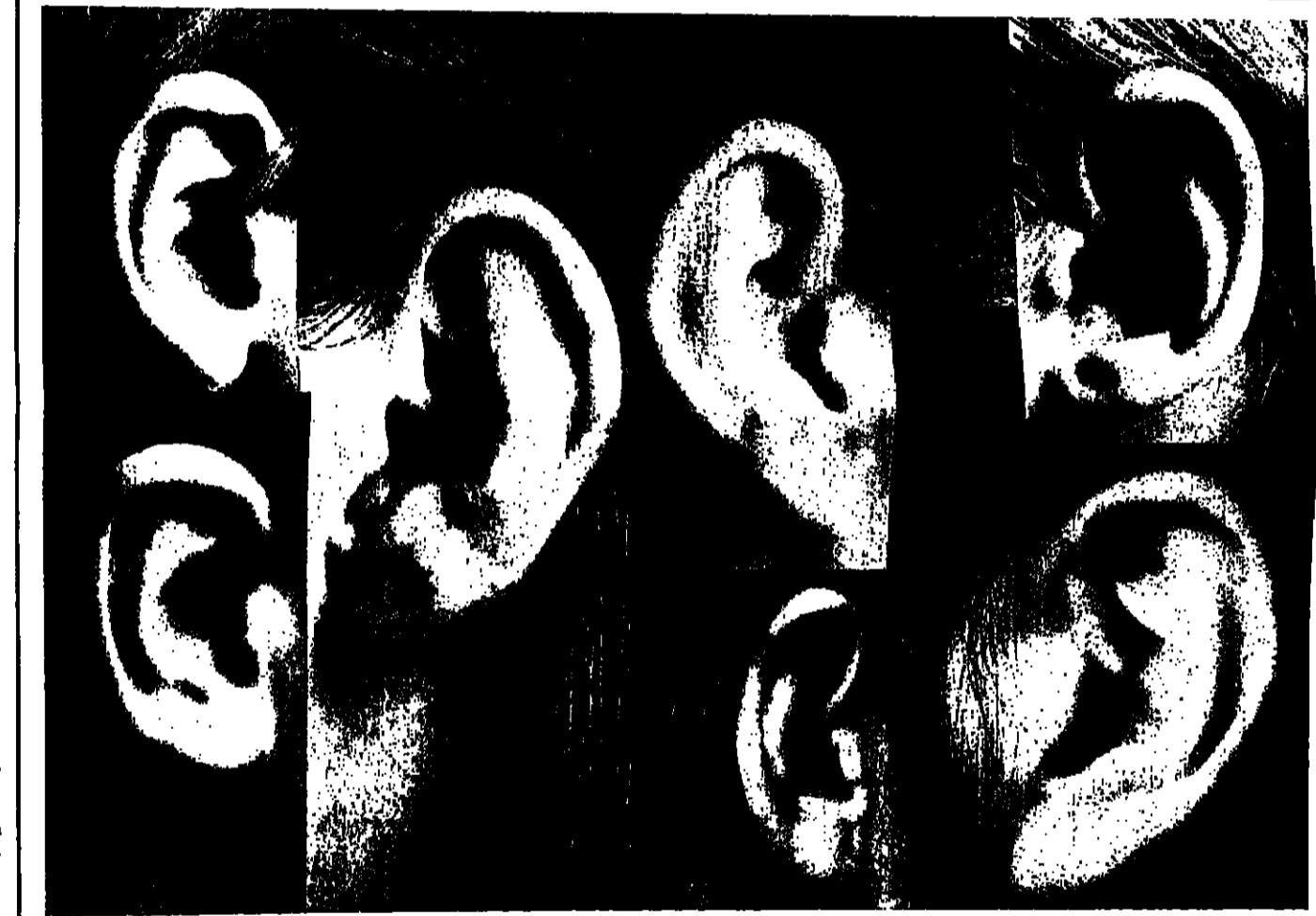
Medical Tribune World Service

**TORONTO**—A commission to review requirements for training neurosurgeons is to be set up soon by the American Board of Neurosurgery, the 64th meeting of the Society of Neurological Surgeons was told.

Dr. W. Kemp Clark of Dallas, Tex., chairman of an A.A.N.S. manpower study, said there will soon be 3,000 neurosurgeons in the United States, including 600

now in training. He noted that most members of the specialty are in their early and late 30s, with very few in the upper age ranges. Based on the present birth rate, Dr. Clark said, there will be one neurosurgeon per 50,000 persons by 1985.

His remarks came after Dr. William Sweet of Boston said a recent study shows that one neurosurgeon per 160,000 persons leaves no large unmet need.



## All Ears!

**CONTRAINdications:** This drug is contraindicated in tuberculous, fungal or viral lesions (herpes simplex, varicella and varicella). It is also contraindicated in those individuals who have shown hypersensitivity to any of its components.

**WARNINGS:** Anecdotal and clinical medical literature indicate an increase in the incidence of sensitivity to neomycin. The possibility of such a reaction should be borne in mind.

**PRECAUTIONS:** As with other antibiotic preparations, prolonged use may result in the overgrowth of non-susceptible organisms. Appropriate measures should be taken if this occurs. Treatment should not be continued longer than 10 days.

**SUPPLIED:** Bottles of 10 cc. and 6 cc. with sterile droppers.

- Broad antibacterial action against susceptible strains of organisms in otitis externa
- Effective concentration of hydrocortisone diminishes edema, itching and pain
- Low pH for acidification
- Convenient 10 cc. size of Cortisporin Otic Drops enables patient to complete a full treatment regimen economically (costs patient about half as much as another leading brand, according to Drug Topics Redbook)

**\*INDICATIONS:** Based on a review of this drug by the National Academy of Sciences—National Research Council and/or other information, FDA has classified the indications as follows:

"Possibly" effective: For the treatment of external otitis either due to or complicated by bacterial infection caused by organisms susceptible to polymyxin B sulfate or neomycin sulfate. It is also valuable in conjunction with systemic therapy in infections of mastoidectomy and fenestration cavities.

Final classification of the less-than-effective indications requires further investigation.

Each oz. contains: Aerospin® brand Polymyxin B Sulfate 10,000 Units; neomycin sulfate 5 mg. (equivalent to 3.5 mg. neomycin base); hydrocortisone 10 mg. (1%). The vehicle contains the inactive ingredients: cetyl alcohol, propylene glycol, polyisobutylene, purified water and thimerosal (preservative) 0.01%.

Complete literature available on request from Professional Services Dept. P.M.C.

 Burroughs Wellcome Co.  
Research Triangle Park  
North Carolina 27708



## When cardiac complaints occur in the absence of organic findings, underlying anxiety may be one factor

### The influence of anxiety on heart function

Excessive anxiety is one of a combination of factors that may trigger a series of maladaptive functional reactions which can generate further anxiety. Often involved in this vicious circle are some cardiac arrhythmias such as paroxysmal supraventricular tachycardia and premature systoles. Since these symptoms resemble those associated with actual organic disease, the overanxious patient needs reassurance that they have no organic basis and that reduction of excessive anxiety and emotional overreaction would be medically beneficial.

### The benefits of antianxiety therapy

Antianxiety medication, when used to complement counseling and reassurance, should be both effective and comparatively free from undesirable side

effects. Extensive clinical experience for more than 13 years has demonstrated that Librium fulfills these requirements with a high degree of consistency. Because of its wide margin of safety, Librium may generally be administered for extended periods, at the physician's discretion, without diminution of effect or need for increase in dosage. (See summary of product information.) If cardiovascular drugs are necessary, Librium is used concomitantly whenever anxiety is a clinically significant factor. (See Precautions.) Librium should be discontinued when anxiety has been reduced to appropriate levels.

### For relief of excessive anxiety and related cardiac dysfunction

adjunctive  
**Librium<sup>®</sup> 10 mg**  
 (chlordiazepoxide HCl)  
 1 or 2 capsules t.i.d./q.i.d.



Roche Laboratories  
 Division of Hoffmann-La Roche Inc  
 Nutley, N.J. 07110

Before prescribing, please consult complete product information, a summary of which follows:

**Indications:** Relief of anxiety and tension occurring alone or accompanying various disease states.

**Contraindications:** Patients with known hypersensitivity to the drug.

**Warnings:** Caution patients about possible combined effects with alcohol and other CNS depressants. As with all CNS-acting drugs, caution patients against hazardous occupations requiring complete mental alertness (e.g., operating machinery, driving). Though physical and psychological dependence have rarely been reported on recommended doses, use caution in administering to addiction-prone individuals or those who might increase dosage; withdrawal symptoms (including convulsions), following discontinuation of the drug and similar to those seen with barbiturates, have been reported. Use of any drug in pregnancy, lactation, or in women of childbearing age requires that its potential benefit be weighed against its possible hazards.

**Precautions:** In the elderly and debilitated, and in children over six, limit to smallest effective dosage (initially 10 mg or less per day) to preclude ataxia or oversedation, increasing gradually as needed and tolerated. Not recommended in children under six. Though generally not recommended, if combination therapy with other psychotropics seems indicated, carefully consider individual pharmacologic effects, particularly in use of potentiating drugs such as MAO inhibitors and phenothiazines. Observe usual precautions in presence of impaired renal or hepatic function. Paradoxical reactions (e.g., excitement, stimulation and acute rage) have been reported in psychiatric patients and hyperactive aggressive children. Employ usual precautions in treatment of anxiety states with evidence of impending depression; suicidal tendencies may be present and protective measures necessary. Variable effects on blood coagulation have been reported very rarely in patients receiving the drug and oral anticoagulants; causal relationship has not been established clinically.

**Adverse Reactions:** Drowsiness, ataxia and confusion may occur, especially in the elderly and debilitated. These are reversible in most instances by proper dosage adjustment, but are also occasionally observed at the lower dosage ranges. In a few instances syncope has been reported. Also encountered are isolated instances of skin eruptions, edema, minor menstrual irregularities, nausea and constipation, extrapyramidal symptoms, increased and decreased libido—all infrequent and generally controlled with dosage reduction; changes in EEG patterns (low-voltage fast activity) may appear during and after treatment; blood dyscrasias (including agranulocytosis), jaundice and hepatic dysfunction have been reported occasionally, making periodic blood counts and liver function tests advisable during protracted therapy.

**Supplied:** Librium<sup>®</sup> Capsules containing 5 mg, 10 mg or 25 mg chlordiazepoxide HCl. Libritabs<sup>®</sup> Tablets containing 5 mg, 10 mg or 25 mg chlordiazepoxide.

The Only Independent Medical Newspaper in the U.S.

# Medical Tribune

## and Medical News

Published by Medical Tribune, Inc.

### Advisory Board

JOHN ADRIANI, M.D. • JULES H. MASSERMAN, M.D. • ROBERT A. CHASE, M.D.  
 ARTHUR M. MASTER, M.D. • RENE J. DUBOS, PH.D. • ALTON OCHSNER, M.D.  
 BERNARD LOWN, M.D. • LEO G. RIGLER, M.D. • ALBERT B. SABIN, M.D.

ARTHUR M. SACKLER, M.D.  
 International Publisher

WILLIAM F. B. O'DONNELL  
 General Manager

RICHARD S. GUBNER, M.D.  
 Associate Editor

NATHAN HORWITZ  
 News Editor

WILLIAM PRIFTIS  
 Layout Editor

880 Third Avenue, New York, N.Y. 10022 • Telephone: 421-4000  
 Circulation audited by Business Publications Audit of Circulation, Inc.



"He says he makes house calls, and he'll be over as soon as he gets his horse back from the blacksmith."

© 1973 Medical Tribune

**HERE** Pleural effusion



Wherever it hurts,  
 Empirin Compound with  
 Codeine usually provides  
 the relief needed.

**HERE** Biliary calculi



In general, only pain so severe  
 that it requires morphine is  
 beyond the scope of  
 Empirin Compound with Codeine.

**prescribing convenience:**  
 Up to 5 refills in 6 months,  
 at your discretion (unless  
 restricted by state law); by  
 telephone order in many states.

Empirin Compound with  
 Codeine No. 3, codeine  
 phosphate\* 32.4 mg. (gr. 1/2);  
 No. 4, codeine phosphate\*  
 64.8 mg. (gr. 1). Warning—  
 may be habit-forming. Each  
 tablet also contains: aspirin  
 gr. 3 1/2, phenacetin gr. 2 1/2,  
 caffeine gr. 1/2.

**Burroughs Wellcome Co.**  
 Research Triangle Park  
 North Carolina 27709

# WHEREVER IT HURTS

**HERE**  
 Osteoarthritis



# EMPIRIN® COMPOUND CODEINE

#3, codeine phosphate\* (32.4 mg.) gr. 1/2  
 #4, codeine phosphate\* (64.8 mg.) gr. 1

A recent report in this newspaper noted that Sir MacFarlane Burnet, the 73-year-old Nobel Laureate, carries a particular medical card in his pocket. It is unlike the usual ones stating, for example, that the bearer is a diabetic. Sir MacFarlane's card reads: "I request that, in view of my age, any prolonged unconsciousness, whether due to accident, heart attack, or stroke, should be allowed to take its course without benefit of an intensive care or resuscitation ward."

Sir MacFarlane observes that modern medical science has become too successful in its ability to prolong life. He adds, "Once I reach the stage of pre-death, all I ask is that I go on to the end with as much dignity and as little pain as possible. Death in the old should be accepted as something always inevitable and sometimes positively desirable. Physicians should not compel old people to die more than once."

It is, of course, true that with modern resuscitative equipment and various supportive measures life can be prolonged or, perhaps more accurately, death can be ward off almost indefinitely, depending on the criteria of life and death. In a simpler past, and in the memory of many physicians, neither the equipment nor the measures were available, and death ensued in due course without any great ability to prevent or delay it.

And not so long ago most deaths occurred at home, rather than in the hospital, because the attending physician had judged hospitalization would be of no

value. Aside from sudden death, most deaths now occur in the hospital. It is estimated that 80 per cent of Americans die within hospitals or other institutions. In a survey of all deaths during a three-week period in an area in Wales, Dr. W. Dewi Rees reported that of 50 people who died, 13 died at home, 26 in a general-practitioner hospital, four in a chronic sick unit, and seven in other hospitals. So, even here, only 26 per cent died at home.

It is with these developments—the availability of potent death-delaying measures, yet the final exit of most people occurring within the hospital—that physicians are being compelled to assess their roles as going beyond that of physicians or, in the dictionary definition, "persons skilled in the art of healing." Within hospitals and medical schools attempts are currently being made to direct attention to the dying. In the past, physicians have been reluctant to do so, often withdrawing from the dying because the experience has been personally too painful, too disturbing.

When the great jurist Oliver Wendell Holmes was 94 years old, he addressed the American people on radio. It was shortly before his death. His final words quoted Thucydides: "Death plucks at my ears and says, 'Live, I am coming.' " Perhaps that means just what Sir MacFarlane's words do: "When the old reach a stage when they cannot cope for themselves, it is true compassion to bring that intolerable stage of pre-death to an end as soon as possible."

Since the cat is now out of the bag, health professions can no longer play ostrich. What is needed is an appropriate response to educate the public about the hemoglobinopathies and to screen and counsel them in an appropriate setting so that misunderstandings will be minimized.

The Metropolitan Seattle Sickle Cell Program is attempting to do just that, and it is questionable whether the 47 families studied by Dr. Hampton were counseled by counselors from the program. Prior to the institution of the Metropolitan Sickle Cell Program last year, however, the problem of confusing sickle cell trait with sickle cell anemia in this community was common. This related to the fact that people who were tested were not counseled or were poorly counseled by the other screening groups in the community.

MAX C. BADER, M.D., M.P.H.  
 Central District Health Officer  
 Seattle, Wash.

## Dietary Punishment

THE COUNCIL ON Foods and Nutrition has said it forcefully, but we think British House Secretary Robert Carr has said it best. Noting that British prisons will abolish bread-and-water diet punishments, the honorable cabinet minister observed that "dietary punishment is out of place in the 20th century." The A.M.A. Council, reviewing Dr. Atkins' *Diet Revolution* and other low-carbohydrate ketogenic weight reduction regimens in the June 4 issue of *J.A.M.A.*, made much of the misery of persons on such diets. Whether it is bread and water or strict avoidance of carbohydrates, we submit that all diets are dietary punishment.

One of Webster's definitions of punishment is "to deplete in quantity, as food or drink;" another "to control or to establish habits of self-control." No less is this true of the low-fat, low-cholesterol diet

achieved, are difficult to sustain. R.S.G.

that is a sacrifice, which is a voluntary form of punishment. The difference between the low-calorie diet, low carbohydrate or otherwise, and the low-fat, low-cholesterol diet is that one can learn to live with fat- and cholesterol-restricted foods whereas caloric restriction and weight reduction, by whatever method

achieved, are difficult to sustain. R.S.G.

achieved, are difficult to

## New Administration Health Plan to Take Hard Line on MDs

*Continued from page 1*  
suggest that "the crux of the problem, to put it plainly, is that you have to keep the doctors honest and then the system will work."

Dr. Edwards replied, "I don't know that I would use the word 'honest'... I think we have to monitor their use of medical technology, be it surgery or the utilization of other kinds of technology, and be sure it is being used both properly... and under the right circumstances."

Secretary of Health, Education, and Welfare Casper W. Weinberger reported that the new national health insurance proposal, which he hopes to show President Nixon next month, is the result of a "searching re-examination" of the 1972 plan.

"It's far from final yet," he said, observing that two options currently receiving close attention are (1) a combination of employer-mandate coverage and federally funded catastrophic protection and (2) a national plan along the lines of the Federal Employees Health Benefits Program.

"Whatever system survives," Mr. Weinberger said, "any plan will include certain concepts. We will propose a partnership concept involving private insurance companies and public agencies, with the public interest and the Government's responsibility to the public, we hope, protected along the way, every step."

"We will assure that all Americans have access to basic comprehensive health insurance and that lack of sufficient income will not be a barrier to obtaining such coverage."

### Would Reduce Cost Inflation

The plan will also include features "that will halt or reduce medical cost inflation and discourage overuse of health care personnel and facilities," he said, adding that "all intentions would be in vain if we have to sit by and watch the value of benefits provided quickly eroded by more inflation."

Noting the "sobering" fact that Medicare patients are now paying just about as much out-of-pocket costs for their health care as they were paying before the program was introduced, Mr. Weinberger went on:

"We believe that health care financing... should be able to be used as a lever to improve the distribution and supply of health care resources."

"We believe that reimbursement mechanisms must be structured so as to encourage the introduction of new concepts, such as health maintenance organizations, physician extenders, and paraprofessional personnel."

"We believe that reimbursement procedures should create new incentives for more efficiency and better quality—concentrate, in other words, on health maintenance rather than on just treatment."

In response to questions, Mr. Weinberger said the PSRO mechanism would be used to help prevent such abuses as "gang visits" as well as other tactics that have permitted "a very few" physicians to make very large incomes entirely from Medicare and Medicaid patients.

### Reimbursement Mechanism Scored

"We have for too long, and in too many situations, been reimbursing health care providers for the exact amount of their charges, without anything near a critical examination of the necessity, the validity, or the propriety of some of those charges," he said.

Mr. Weinberger said past Government policies were to blame for encouraging—or, at least, not restraining—questionable practices, but a major effort is now under way to "remove any Government stimulus there might be to more health care inflation."

Picking up this theme later, Dr. Edwards said the most important issue is the need to develop a "national health strategy" that will bring the "vast resources" of the public and private sector to bear on the problems of rising costs,

maldistribution, unequal access, and quality of care.

In sharp contrast with the conciliatory tone of his recent address to the American Medical Association meeting, Dr. Edwards enumerated a number of physician-related problems that the national health strategy might solve—and some that it could not.

One problem that will be addressed is the "so-called doctor shortage"—which is really a problem of distribution. The real problem, Dr. Edwards said, is that "there are not enough doctors providing primary care, while the number of specialists—general surgeons and the like—appears to be greater than necessary and even, for that matter, increasing."

"In addition, we are not making effective—I emphasize 'effective'—use of allied health care professionals," he said.

Some years ago the threat of a doctor shortage resulted in increased Federal aid for medical education. Medical school enrollment is up, and by the latter half of this decade between 9,000 and 10,000 new physicians will be graduating each year.

This increase will not, in and of itself, correct the shortage of primary care physicians or the oversupply of specialists, Dr. Edwards said, noting that we may end up with a doctor surplus.

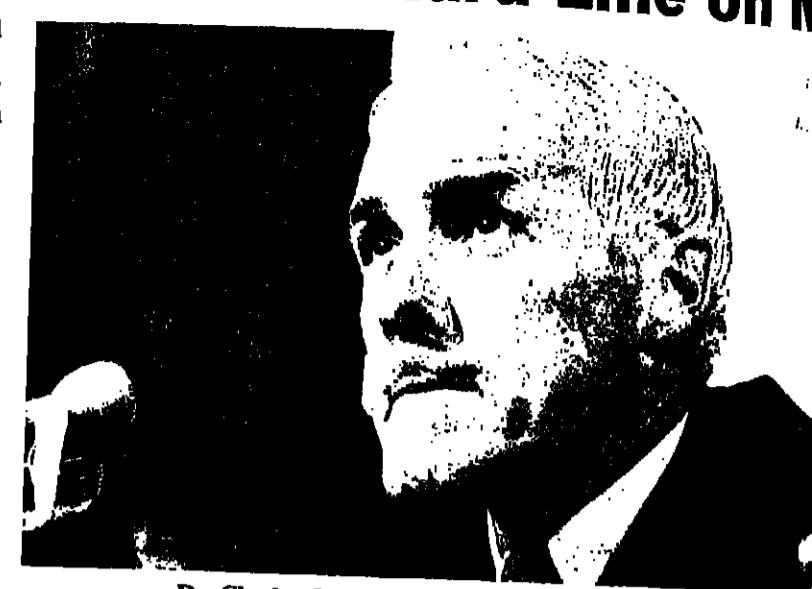
"If that were to happen," he said, "you might assume that physicians' fees would fall under the influence of normal supply-and-demand factors."

"I think the reverse is likely to happen. I think it already has happened. An excess of physicians, like an excess of hospital beds, tends to increase demand and certainly not lower it."

The we could be looking toward even greater inflation in the cost of health care as surplus doctors try to generate a certain amount of demand for their services," he said.

Dr. Edwards, a former surgeon, cited some "astounding" statistical comparisons between the United States and England.

"The number of full-time surgeons in the United States per 1,000 population is about 37.6 versus 20.8 in England," he said. "The number of operations per 1,000



Dr. Charles C. Edwards, Assistant Secretary for Health. (Wide World)

population in the United States in 1969 was 73.2. In England it was 46.4.

The number of tonsillectomies per 1,000 population in the United States was 6.3 compared with 3.3 in the U.K."

Dr. Edwards said that "if we launch a national health insurance system and fail to address this kind of utilization issue, I think we would be failing our responsibility very badly."

One of the problems that the Federal Government cannot handle by itself, Dr. Edwards said, is the misuse of certain drugs, particularly the antibiotics.

Recent studies indicate that as many as 60 per cent of hospitalized patients on antibiotic therapy had no evidence of infection," he said. "Other studies suggest that about 30 per cent of the patients who see a doctor because of the common cold receive a prescription for one of the commonly used antibiotics."

"The problem here is not simply that these drugs will do the patient no real good," he went on. "They can do serious harm by promoting the growth of resistant strains of bacteria and an increased number of superinfections against which conventional antibiotic therapy proves ineffective."

Dr. Edwards said the FDA—of which he is a former commissioner—"provides physicians with accurate and complete information on drug usage, and I think it is improving in this regard."

"But it is plain that many physicians are not making proper use of this information, and if they were, we wouldn't be producing and certifying enough antibiotics each year to supply 50 doses a year for every man, woman, and child in the United States."

### Asked About PSRO

Asked how he felt about A.M.A. opposition to PSRO, Dr. Edwards replied:

"Well, I am not so naive as to think that the American Medical Association is going to fall in lock step and go down the road with us on PSRO without some major biffles. They obviously aren't."

"I think, though, there are sufficient numbers of enlightened physicians in the United States that are beginning to recognize the fact that the medical profession has been practicing over these many years will little or no quality control," he said.

"I think that we are slowly bringing these people into the real world, which recognizes that we all—including those of us in Government—have a hell of a lot of people looking over our shoulder and providing a certain amount of quality control."

## 3 Nonsystemic Contraceptives Show Promise

*Continued from page 1*

moved intact, and no adverse tissue reaction, inflammation, or histologic changes were seen.

Citing clinical trials of such plugging conducted in India, Dr. Erb noted that antifertility efficacy has been achieved for as long as two years even though investigators there used a highly diluted, low-viscosity silicone polymer and simply filled the uterine cavity with this material.

A non-air-entrapning mixer and dispenser devised by the Philadelphia team, he added, permits the use of higher-viscosity polymer systems. The advantages include prevention of intraperitoneal spillage, good conformance to the oviduct lumen, and higher tensile strength.

Dr. Erb said the development of a special obturator injection tip has been another factor in the success of the Philadelphia experiments. The tip's structure facilitates delivery of the polymer into the fallopian tube without significant backflow into the uterine cavity. Also, the tip detaches from the insertion device and remains in the uterus, bonded to the cured oviduct plug.

For use in women, the tips would be fitted with loops or other means of attachment to allow for transcervical removal and thus permit reversibility of the contraceptive sterilization.

Co-authors of the report were Robert H. Davis, Ph.D., and Dr. George A. Kyriazis and Howard Ballin.

A new intrauterine device, consisting of

a soft pouch that is inflated with sterile saline after insertion, has been developed by Dr. Jack M. Futran and Sotiris Kitrakis, of the University of California School of Medicine, San Francisco.

The pouch has no sharp edges or points to irritate the uterus, Dr. Futran pointed out. And the postinsertion inflation means that the device is, in effect, custom-fitted, he said.

A total of 207 women, more than half of them nulliparous, have been fitted with the device for a total of about 600 months of exposure. No pregnancy has occurred with the device in place.

Five of the devices have been expelled and five others removed because of bleeding and/or pain during the trials, which have provided an average follow-up time of three months. Most problems with an IUD become evident within the first three months, Dr. Futran commented.

The investigators emphasized that their results must be considered preliminary because of the relatively small number of patients. They believe, however, that the incidence of expulsions and removals has been "encouragingly low," especially in a study sample that includes many nulliparous women, and they say that patient reaction has been "enthusiastic."

A second adaptation of the intrauterine device—one that delivers very small doses of progesterone—was described by Dr. Antonio Scormegna, of the University of Chicago Pritzker School of Medicine.

This local administration of progeste-

ron in minute amounts induces decidua changes in the endometrial receptor that make it unsuitable for implantation, but the hormone does not have systemic effects, Dr. Scormegna said.

The vehicle used in the clinical trials has been a small, T-shaped device, the so-called Tatum T. It has a low expulsion rate and seldom causes bleeding or pain but is associated with a relatively high pregnancy rate.

Dr. Scormegna called the results so far achieved with the progesterone-releasing device "quite encouraging." No pregnancy has occurred while a device releasing adequate amounts of progesterone was present in the uterine cavity.

The rates of expulsion and cervical displacement, as well as the incidence of removal for medical or personal reasons, were higher than those reported for the regular T device. But Dr. Scormegna noted that repeated removals and insertions probably contributed to this increase.

The experimental device had a useful life of only six months because of gradual hormone release.

Further factors influencing removal were the endo-

metrial biopsies and other procedures required by the study and the prevalence of pelvic inflammatory disease in the study population.

Associated with him in the study were Drs. Theresita Avila, Manuel Luna, Ramona Rao, and W. Paul Dmowski and B. Kulkarni, Ph.D.

Medical Tribune

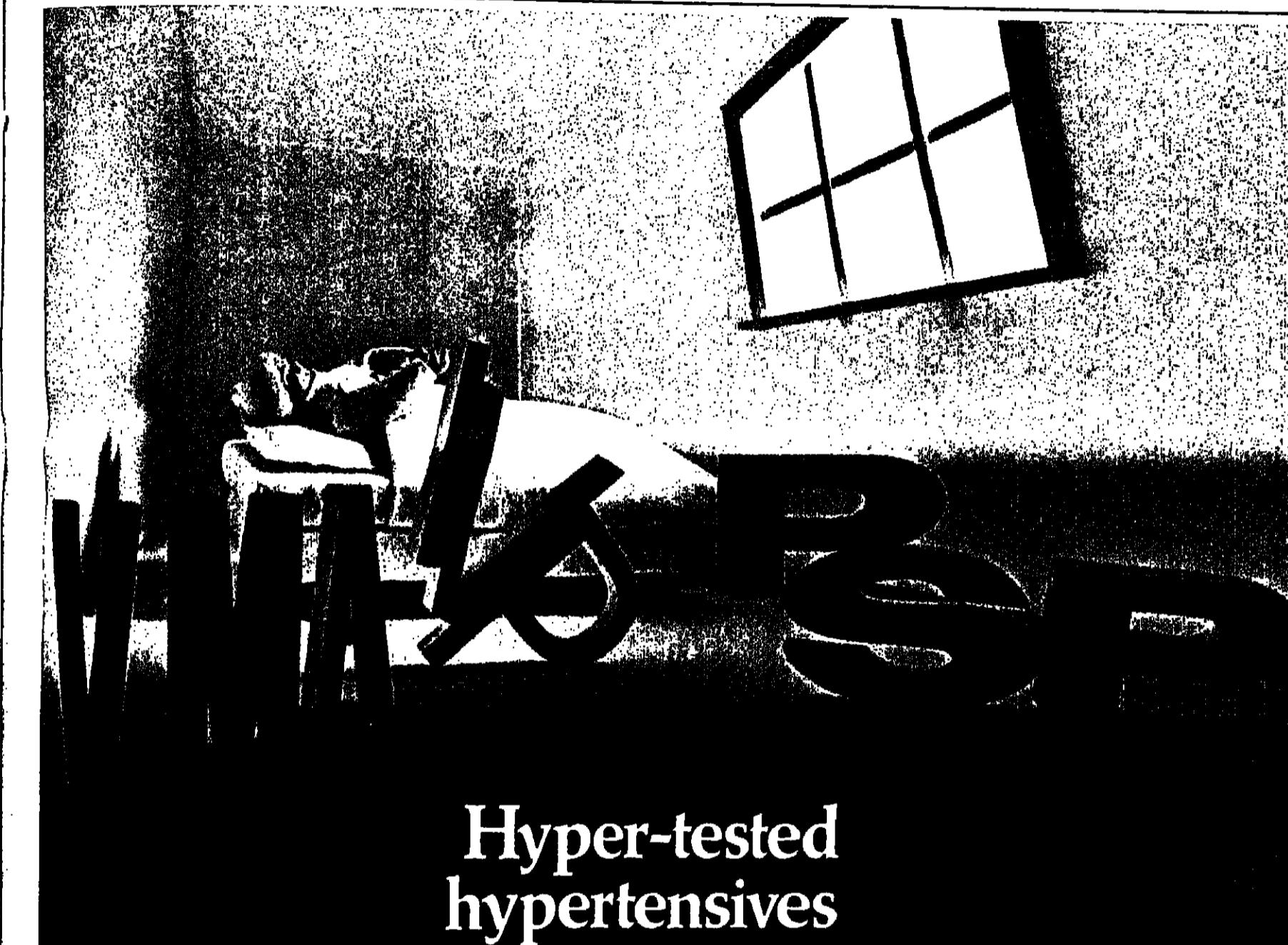
# HYPERTENSION BULLETIN

ACIBA SERVICE

Table of Contents	BP and me	18
	Reports from abroad	19
	Artist of medicine	20
	Mechanics of stress	25
	No-pump dialyzer	26
	Hypertension classics	27

AUGUST 8, 1973

PREPARED BY INTERNATIONAL MEDICAL PRESS



## Hyper-tested hypertensives

THE COMMON PRACTICE of doing the big work-up on patients with mild to moderately severe hypertension is all wrong, says Dr. Marvin Moser, a White Plains, N.Y. cardiologist to whom Manhattan nephrologists often send patients when they can't get blood pressures down to satisfactory levels.

"For the past 15 years, many physicians have been looking mainly for the esoteric, the unusual. They hospitalize the patient, searching for the three to five per cent who have fascinating complications. That may be much more interesting scientifically, but it's not real therapy."

"The real therapy begins with a blood pressure cuff and pills, not with elaborate testing, not with IVPs, VMAs, PSPs and all the rest of the diagnostic alphabet

soup. I'm talking about something as simple as a diuretic three times a week to start, or a diuretic combined with a tranquilizer, rauwolfa, or a barbiturate. Later on, the use of higher dosages and other drugs may be necessary. Not very glamorous. Won't get you published. But it will help you help your patients and may make a dent in the enormous morbidity due to hypertensive disease."

From 1958 to 1971 Dr. Moser, who is the author with Dr. Arthur Goldman of *Hypertensive Vascular Disease*, was physician-in-charge of the hypertension service at Montefiore Hospital and Medical Center in New York City and he still serves the hospital as an associate attending in medicine. He is also chief of cardiology, White Plains Hospital, consultant

continued on page 18

## Hyper-tensioned hypertensives

continued from page 17

in cardiology to the State Department of Health, and Assistant Professor of Clinical Medicine, Albert Einstein College of Medicine.

Dr. Moser consistently treats patients whose repeated blood pressure readings are as low as 140/90, especially if they are under 40 and if there are the risk factors of familial history of early coronary or hypertensive disease. He even treats patients in the 40-60 age group whose pressures are in the 145-150/90 range.

"Let's take a man of 50, with a blood pressure of 150/95 and an uncle who died at 55 of a stroke. I would probably not treat him at once, but would ask him to come back in three to six months. If the

### The real therapy begins with a blood pressure cuff and pills, not with elaborate testing.

pressure were then above 150/90, I would start him on 50 mg. of hydrochlorothiazide three times a week, possibly combined with a rauwolfa preparation, unless he had a history of depression or appeared to be a highly anxious person.

"Now suppose him to be obese. I would get him to lose weight, but I would know that this is only likely to control blood pressure if it is mildly elevated. A lot of physicians, I think, rely too much on that notion in severe cases and neglect treatment. The fact remains, however, that obesity is another risk in heart disease. If his cholesterol were high, I'd try to correct that, and if he smoked two packs of cigarettes a day, I would go after that, too. These are all peripheral to hypertension, but contributors to coronary disease."

Dr. Moser does not see a patient frequently unless his hypertension is both

When Dr. Moser needs to consult about a particularly tricky problem he turns to his American Indian kachina associate for guidance and inspiration.

severe and refractory. And further: "No scare tactics, no telling him his heart may give out or his kidneys may fail or a stroke may result. I think when someone finds out he has high blood pressure he is frightened enough. Everyone knows someone or of someone who had a stroke.

As a member of a joint American Heart Association-American College of Cardiology Self-Assessment Study Group on Hypertension, Dr. Moser is sold on the idea of a "cookbook" approach to treatment of mild to moderately severe hypertension.

"I know that some people object to the 'cookbook' concept. They think they are not individualizing treatment, but our experience pretty conclusively shows that about 60 per cent of all mild to moderately severe hypertensives will respond to thiazide and rauwolfa in medium to full doses—up to 100 mg. of each per day. That's a pretty high percentage. It means that if we screened the whole country and found eight million more such hypertensives, almost two thirds of them could be successfully treated by the so-called 'cookbook' prescription."

For such patients, Dr. Moser's plan of therapy is placid, and persistent. He is going to get that reading down, but he is not going to disrupt the patient's customary way of life in the process.

"I don't believe that all hypertensives, or even 20 per cent of them for that matter, should have elaborate work-ups. I don't think they should be in hospitals. I feel very strongly that the vast majority of hypertensives who are lying in hospitals for work-ups don't belong there."

This does not apply to patients with malignant hypertension, of course. They belong in the hospital for emergency treatment.

"That is the patient with a diastolic of 130-140 plus, with headaches, dizziness,



# 135/85

DR. IRVING H. PAGE, director emeritus of research at the Cleveland Clinic, and the man who led one of two research teams that separately reported the discovery of angiotensin in 1939, took his own blood pressure for years, but now he has given way and has another doctor check him once a year. Dr. Page, at 72, says his pressure holds at about 135/85, partly because of a "reasonably good heredity".

He jogs, plays tennis and, "to the great distress of friends," walks and walks. He began to "monkey around" with low fat and low cholesterol intake in the 1940s, but did not give up alcohol. Smoking he did stop, after 50 years.

He has never been convinced that stress alone can cause high blood pressure (see his article: *Hypertension Bulletin*, June), but sees the issue as irrelevant in his own case, since "things don't really stress me terribly".

①

"By now it's clear that this is a patient who has to be investigated for a possible renovascular lesion, a narrowing of one

or both renal arteries. In a man this age, the most common cause would be an arteriosclerotic plaque that you hope will be in the renal artery, just where it comes off the aorta, so that surgery can be effective. In a young woman, you might find a different kind of lesion—a dysplasia of the media or intima of the renal artery. These lesions are frequently not amenable to surgical repair."

But such situations are rare, and Dr. Moser does not work up patients for renovascular disease unless the disease is progressive, non-responding, or accelerating. However, he believes that every hypertensive under 30 should be worked up, because that is where the yield really is in the search for abnormalities.

The remainder, he treats at once, after giving a basic, baseline examination: An ECG indicates possible heart involvement as a target organ. A simple urine, if negative, rules out renal parenchymal disease. PSP tests and creatinine clearances are unnecessary.

"These tests are nice from an academic point of view, and I think interns and residents ought to learn about them, but from a practical standpoint we don't need them. A BUN and a urine will tell you as much as you really need to know to begin treatment of all but the malignant or severely progressive hypertensives. We have a serious public health problem with untreated hypertensives and we will not begin to solve it unless we simplify our approach!"

Even where there is kidney damage, as evidenced, say, by a BUN of 25 to 35 mg. per cent or a creatinine of 1.5 to 2.5 mg. per cent, there is no reason to withhold antihypertensives.

"I have seen a fair number of people both in private practice and at Montefiore who started out with evidence of kidney

**A BUN and a urine will tell you as much as you really need to know to begin treatment of all but the severely progressive.**

insufficiency—BUNs of 30 and 40 mg. per cent and creatinines of 2, 3 or even 4 mg. per cent—whose function has remained stable for five or more years following effective antihypertensive treatment.

Dr. Moser recalls only two patients in whom a creatinine of over 4 mg. per cent came down toward normal after treatment, and adds that there are some investigators who have reported more frequent improvement in kidney function.

He explains this phenomenon in terms of the effect of hypertension on target organs. High blood pressure, he believes, can be regarded as simply a mechanical burden on the heart as a pump and the blood vessels of the kidney and brain, whatever the cause of the elevation.

"If you can take away this mechanical insult, this added blood pressure, you

have eliminated a major factor in the etiology of vessel disease in these major target organs."

The fallacy persists, he says, that as you lower blood pressure, you reduce blood flow to the kidneys, decrease glomerular filtration and cause further kidney damage.

"This is true—temporarily, especially with drugs like guanethidine, which act by dilating the veins, reducing venous re-

**I don't believe that all hypertensives, or even 20 per cent of them, should be in hospitals.**

turn to the heart, thus decreasing cardiac output. When cardiac output falls, so does the flow to the kidney. With such drugs you frequently see rising BUN levels, indicating less blood flow to the kidney as the pressure comes down.

"House officers are fond of making charts showing the patient's blood pressure coming down and the kidney function getting worse.

"What are you doing to this poor fellow?" they would ask us during rounds. The answer to that is, that if you drop the blood pressure of patients with renal insufficiency carefully, and if you persist in treating them, the BUNs will first rise and will then come back down to pretreatment levels and stay there. After a month to six weeks, as blood pressure comes under control, the BUN and creatine levels will come back down."

By far the greater danger in such patients is inadequate treatment, Dr. Moser says. He cites a man in his early 50s with diastolics of 140-150-160, heart failure, a creatinine of 6 mg. per cent, and retinal hemorrhages.

"You knew very well that by lowering his pressure you were not going to prolong his life indefinitely, because his kidney function was too far gone. Treatment proved to be more than just palliative, however. This man has been on therapy about a year and a half. His creatinine is still six to seven, his BUN is 90 to 100. He is obviously losing ground in renal function, but he has had 18 months of pretty good health. His fundi are free of hemorrhage; he's out of heart failure; and he's going about his business.

"I like to think that, if he had been treated five years ago, this never would have happened at all. The sad part of the story is that he had been seeing a doctor all along, but unfortunately he had been given totally inadequate therapy.

"This case is characteristic of something you see frequently: the patient whose doctor is aware that the hypertension ought to be treated and therefore uses a drug, or even two or three drugs, in inadequate dosage. The doctor goes to bed at night feeling very comfortable. He is treating his hypertensive patient, but is not treating him to end-point."

## reports from abroad

HELSINKI—The results of a serum lipid and lipoprotein study at the University of Helsinki by Drs. Isko Nikkila and Antti Aro suggest a familial trait of hyperlipidemia in one-third of families in which one member had suffered premature coronary heart disease.

Of the 101 families examined, hyperlipoproteinemia was prominent in 33 and only nine of these had a single-type disease. Familial hypercholesterolemia (type-IIa) was found in six families. Abnormal lipoprotein phenotypes coexisted in 24 families, with half the members affected. Phenotype IIb was six times more common in the first-degree relatives of myocardial infarction survivors than in a control population.

JERUSALEM—Profound salt wastage found in seven children with very high plasma-renin activity and normal or high plasma-aldosterone levels may be explained, said Dr. A. Rösler, Hadassah University Hospital, by the non-responsiveness of the renal tubule to aldosterone. The patients responded to heavy dietary supplements of salt.



DUNDEE, SCOTLAND—Patients with chronic renal failure on maintenance hemodialysis "may be subject to a dangerous combination of symptomless duodenal ulceration associated with severe, prolonged hypersecretion of acid" that may be a consequence of the renal disease or its treatment by hemodialysis, report Drs. A. M. M. Shepherd, W. K. Stewart, and K. G. Wormsley, of Maryfield Hospital.

They found very high overnight and basal gastric acid secretion associated with duodenal ulcer in 53 per cent of 15 patients with end-stage chronic renal failure. They suggest that all such patients on hemodialysis be monitored for spontaneous gastric hypersecretion, so that proper therapy can be initiated.



# Two ways to treat moderate hypertension and why...



## why Ser-Ap-Es®

reserpine 0.1 mg  
hydralazine hydrochloride 25 mg  
hydrochlorothiazide 15 mg

because only Ser-Ap-Es adds hydralazine to rauwolfia-thiazide



Ser-Ap-Es does more than control blood pressure in moderate hypertension—it's a therapeutic approach that considers the whole patient. And adding hydralazine to rauwolfia-thiazide

usually permits lower dosage of each component than if prescribed alone.

If there is slight renal impairment, hydralazine helps maintain or increase renal blood flow.

If the patient is stress reactive, the reserpine component should have a calming effect.

If the patient is uncooperative, Ser-Ap-Es may be a help because it contains all the medication many patients need in a single tablet.

Ser-Ap-Es should be used with caution in patients with advanced renal damage and cerebrovascular accidents. It should be discontinued at the first sign of mental depression.

## why Esimil®

guanethidine monosulfate 10 mg  
hydrochlorothiazide 25 mg

because Esimil offers the control-with-convenience so many hypertensives need



Esimil, an equally valuable yet different approach to moderate hypertension, makes sense for many patients because it anticipates future problems while helping to solve present ones.

If the patient is free of organ damage, Esimil may help keep her that way because it provides guanethidine, perhaps the most effective antihypertensive available. And effective lowering of blood pressure takes pressure off target organs.

If the patient forgets things, Esimil may make it easier to remember with once-a-day dosage, feasible in most cases.

Postural hypotension may occur with the use of Esimil, particularly while the drug is being introduced. Like all antihypertensives, Esimil should be given with caution in the presence of severe coronary insufficiency or recent myocardial infarction.

early, effective control of hypertension can save lives

# Ser-Ap-Es®

reserpine 0.1 mg  
hydrochlorothiazide 25 mg  
hydrochlorothiazide 15 mg

# Esimil®

guanethidine monosulfate 10 mg  
hydrochlorothiazide 25 mg

# Ser-Ap-Es®

reserpine 0.1 mg  
hydrochlorothiazide 25 mg  
hydrochlorothiazide 15 mg

**INDICATIONS**  
Based on a review of this drug by the National Academy of Sciences-National Research Council and/or other information, FDA has classified the indications as follows:

Effective: Hypertension. (See box warning.)

**WARNING**  
This fixed combination drug is not indicated for initial therapy of hypertension. Hypertension requires therapy titrated to the individual. This fixed combination drug is less convenient in patient management. The treatment of hypertension is not static, but must be reevaluated as conditions in each patient warrant.

## CONTRAINDICATIONS

**Reserpine**  
Known hypersensitivity, mental depression (especially with suicidal tendencies), active peptic ulcer disease, colitis, and patients receiving electroconvulsive therapy.

**Hydralazine**  
Hypersensitivity to hydralazine; coronary artery disease; and mitral valvular rheumatic heart disease.

**Hydrochlorothiazide**  
Anuria; hypersensitivity to this or other sulfonamide-derived drugs. The routine use of diuretics in an otherwise healthy pregnant woman with or without mild edema is contraindicated and potentially hazardous.

**WARNINGS**  
**Reserpine**  
Extreme caution should be exercised in treating patients with a history of mental depression. Discontinue the drug at first sign of dependency. Discontinuing reserpine after long-term use of this drug may persist for several months after drug withdrawal and may be severe enough to result in suicide.

**Hydralazine**  
Chronic administration of doses over 400 mg per day may produce in a few patients an arthritic-like acute systemic lupus erythematosus. This syndrome may also occur at lower doses. Hyperthyroidism and signs usually regress when the drug is discontinued. Long-term treatment with steroids may be necessary to control the syndrome. Complete blood counts, and antinuclear antibody titer determinations are indicated before and periodically during therapy.

**Hydrochlorothiazide**  
Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. In patients with severe renal disease, the drug may develop in patients with impaired renal function. Thiazides should be used with caution in patients with impaired hepatic function or progressive liver disease, since alterations of fluid and electrolyte balance may be precipitated. Thiazides may be additive or potentiative in the action of other antihypertensive drugs. Potentiation occurs with ganglion and peripheral adrenergic blocking agents.

**Sensitivity reactions**  
Sensitivity reactions may occur in patients with a history of allergy or bronchial asthma.

**The possibility of exacerbation or activation of systemic lupus erythematosus has been reported.**

**Use in Pregnancy**  
**Reserpine**  
The safety of reserpine for use during pregnancy or lactation has not been established; therefore, the drug should not be used in pregnant patients or in women of childbearing potential only when, in the judgment of the physician, the potential benefit to the welfare of the patient, increased maternal or neonatal tract secretions, nasal congestion, coryza, and anuria, may occur in neonates and breast-fed infants of reserpine-treated mothers since this drug crosses the placental barrier and appears in breast milk.

**Hydrochlorothiazide**  
Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. In patients with severe renal disease, the drug may develop in patients with impaired renal function. Thiazides should be used with caution in patients with impaired hepatic function or progressive liver disease, since alterations of fluid and electrolyte balance may be precipitated. Thiazides may decrease serum PBI levels without signs of thyroid disturbance.

**ADVERSE REACTIONS**  
**Reserpine**  
Rauwolfia preparations have caused gastrointestinal reactions, including hypertension, nausea, vomiting, anorexia, constipation, diarrhea, and cardiovascular reactions including angina-like symptoms, tachycardia, bradycardia, central nervous system reactions, including drowsiness, depression, nervousness, paroxysmal hypertension, nightmares, rare parkinsonian symptoms and psychomotor tract symptoms, and CNS sensitization. These reactions are often associated with ganglion and peripheral adrenergic blocking agents.

**Sensitivity reactions**  
Sensitivity reactions may occur in patients with a history of allergy or bronchial asthma.

**The possibility of exacerbation or activation of systemic lupus erythematosus has been reported.**

**Use in Pregnancy**  
**Reserpine**  
The safety of reserpine for use during pregnancy or lactation has not been established; therefore, the drug should not be used in pregnant patients or in women of childbearing potential only when, in the judgment of the physician, the potential benefit to the welfare of the patient, increased maternal or neonatal tract secretions, nasal congestion, coryza, and anuria, may occur in neonates and breast-fed infants of reserpine-treated mothers since this drug crosses the placental barrier and appears in breast milk.

**Hydrochlorothiazide**  
Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. In patients with severe renal disease, the drug may develop in patients with impaired renal function. Thiazides should be used with caution in patients with impaired hepatic function or progressive liver disease, since alterations of fluid and electrolyte balance may be precipitated. Thiazides may be additive or potentiative in the action of other antihypertensive drugs. Potentiation occurs with ganglion and peripheral adrenergic blocking agents.

**Sensitivity reactions**  
Sensitivity reactions may occur in patients with a history of allergy or bronchial asthma.

**The possibility of exacerbation or activation of systemic lupus erythematosus has been reported.**

**Use in Pregnancy**  
**Reserpine**  
Since reserpine increases gastrointestinal motility and secretions, it should be used cautiously in patients with a history of peptic ulcer, ulcerative colitis, or gallstones (biliary colic may be precipitated).

**Caution** should be exercised when treating hypertensive patients with reserpine, since they adjust poorly to lowered blood pressure levels.

**Reserpine**  
Since reserpine increases gastrointestinal motility and secretions, it should be used cautiously in patients with a history of peptic ulcer, ulcerative colitis, or gallstones (biliary colic may be precipitated).

**Caution** should be exercised when treating hypertensive patients with reserpine, since they adjust poorly to lowered blood pressure levels.

**Reserpine**  
Since reserpine increases gastrointestinal motility and secretions, it should be used cautiously in patients with a history of peptic ulcer, ulcerative colitis, or gallstones (biliary colic may be precipitated).

**Preoperative withdrawal** of reserpine does not ensure that circulatory instability will not occur, and the physician should be aware of the patient's history and consider this in the overall management, since hypertension may occur in patients receiving rauwolfia preparations. Metaraminol (norepinephrine) have been employed to treat adverse vasoconstrictive effects.

**Hydralazine**  
Myocardial stimulation produced by hydralazine and/or digitalis and ECG changes of the myocardial ischemia. This drug has been implicated in the production of myocardial ischemia, and must, therefore, be used with caution in patients with suspected coronary artery disease.

## Two ways to treat moderate hypertension



The "hypodynamic" circulation caused by hydralazine may accentuate specific cardiovascular inadecuaciones. An example is that hydralazine may increase the blood pressure in patients with mitral valvular disease. The drug may produce the pressor responses to epinephrine. Postural hypotension may result from hydralazine but is less than with ganglion blocking agents. Use is indicated in patients with cerebral vascular accidents.

In hypertensive patients with normal kidneys, who are treated with hydralazine, there is evidence of a decrease in glomerular flow and a maintenance of glomerular filtration rate. Instances improved renal function has been noted where control values were below normal prior to hydralazine therapy. However, as with any antihypertensive agent, hydralazine should be used with caution in patients with advanced renal damage.

**Peripheral neuropathy**, evidenced by paresthesias, numbness, and loss of reflexes, has been observed. Published evidence suggests a sympathetic effect and the addition of pyridoxine to the regimen if symptoms develop.

**Eye dysrhythmias**, consisting of reduction in hamartoma, papilledema, and purpura, have been reported. Such abnormalities develop, discontinuing therapy. Periodic blood counts are advised during therapy.

**CONTRAINDICATIONS**

**Reserpine**  
Known hypersensitivity, mental depression (especially with suicidal tendencies), active peptic ulcer disease, colitis, and patients receiving electroconvulsive therapy.

**Hydralazine**  
Hypersensitivity to hydralazine; coronary artery disease; and mitral valvular rheumatic heart disease.

**Hydrochlorothiazide**  
Anuria; hypersensitivity to this or other sulfonamide-derived drugs. The routine use of diuretics in an otherwise healthy pregnant woman with or without mild edema is contraindicated and potentially hazardous.

**WARNINGS**

**Reserpine**  
Extreme caution should be exercised in treating patients with a history of mental depression. Discontinue the drug at first sign of dependency.

**Hydralazine**  
Chronic administration of doses over 400 mg per day may produce in a few patients an arthritic-like acute systemic lupus erythematosus. This syndrome may also occur at lower doses. Hyperthyroidism and signs usually regress when the drug is discontinued. Long-term treatment with steroids may be necessary to control the syndrome. Complete blood counts, and antinuclear antibody titer determinations are indicated before and periodically during therapy.

**Hydrochlorothiazide**  
Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. In patients with severe renal disease, the drug may develop in patients with impaired renal function. Thiazides should be used with caution in patients with impaired hepatic function or progressive liver disease, since alterations of fluid and electrolyte balance may be precipitated. Thiazides may be additive or potentiative in the action of other antihypertensive drugs. Potentiation occurs with ganglion and peripheral adrenergic blocking agents.

**Sensitivity reactions**  
Sensitivity reactions may occur in patients with a history of allergy or bronchial asthma.

**The possibility of exacerbation or activation of systemic lupus erythematosus has been reported.**

**Use in Pregnancy**  
**Reserpine**  
The safety of reserpine for use during pregnancy or lactation has not been established; therefore, the drug should not be used in pregnant patients or in women of childbearing potential only when, in the judgment of the physician, the potential benefit to the welfare of the patient, increased maternal or neonatal tract secretions, nasal congestion, coryza, and anuria, may occur in neonates and breast-fed infants of reserpine-treated mothers since this drug crosses the placental barrier and appears in breast milk.

**Hydralazine**  
Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. In patients with severe renal disease, the drug may develop in patients with impaired renal function. Thiazides should be used with caution in patients with impaired hepatic function or progressive liver disease, since alterations of fluid and electrolyte balance may be precipitated. Thiazides may be additive or potentiative in the action of other antihypertensive drugs. Potentiation occurs with ganglion and peripheral adrenergic blocking agents.

**Sensitivity reactions**  
Sensitivity reactions may occur in patients with a history of allergy or bronchial asthma.

**The possibility of exacerbation or activation of systemic lupus erythematosus has been reported.**

**Use in Pregnancy**  
**Reserpine**  
The safety of reserpine for use during pregnancy or lactation has not been established; therefore, the drug should not be used in pregnant patients or in women of childbearing potential only when, in the judgment of the physician, the potential benefit to the welfare of the patient, increased maternal or neonatal tract secretions, nasal congestion, coryza, and anuria, may occur in neonates and breast-fed infants of reserpine-treated mothers since this drug crosses the placental barrier and appears in breast milk.

**Hydrochlorothiazide**  
Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. In patients with severe renal disease, the drug may develop in patients with impaired renal function. Thiazides should be used with caution in patients with impaired hepatic function or progressive liver disease, since alterations of fluid and electrolyte balance may be precipitated. Thiazides may be additive or potentiative in the action of other antihypertensive drugs. Potentiation occurs with ganglion and peripheral adrenergic blocking agents.

**Sensitivity reactions**  
Sensitivity reactions may occur in patients with a history of allergy or bronchial asthma.

**The possibility of exacerbation or activation of systemic lupus erythematosus has been reported.**

**Use in Pregnancy**  
**Reserpine**  
The safety of reserpine for use during pregnancy or lactation has not been established; therefore, the drug should not be used in pregnant patients or in women of childbearing potential only when, in the judgment of the physician, the potential benefit to the welfare of the patient, increased maternal or neonatal tract secretions, nasal congestion, coryza, and anuria, may occur in neonates and breast-fed infants of reserpine-treated mothers since this drug crosses the placental barrier and appears in breast milk.

**Hydralazine**  
Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. In patients with severe renal disease, the drug may develop in patients with impaired renal function. Thiazides should be used with caution in patients with impaired hepatic function or progressive liver disease, since alterations of fluid and electrolyte balance may be precipitated. Thiazides may be additive or potentiative in the action of other antihypertensive drugs. Potentiation occurs with ganglion and peripheral adrenergic blocking agents.

**Sensitivity reactions**  
Sensitivity reactions may occur in patients with a history of allergy or bronchial asthma.

**The possibility of exacerbation or activation of systemic lupus erythematosus has been reported.**

**Use in Pregnancy**  
**Reserpine**  
The safety of reserpine for use during pregnancy or lactation has not been established; therefore, the drug should not be used in pregnant patients or in women of childbearing potential only when, in the judgment of the physician, the potential benefit to the welfare of the patient, increased maternal or neonatal tract secretions, nasal congestion, coryza, and anuria, may occur in neonates and breast-fed infants of reserpine-treated mothers since this drug crosses the placental barrier and appears in breast milk.

**Hydralazine**  
Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. In patients with severe renal disease, the drug may develop in patients with impaired renal function. Thiazides should be used with caution in patients with impaired hepatic function or progressive liver disease, since alterations of fluid and electrolyte balance may be precipitated. Thiazides may be additive or potentiative in the action of other antihypertensive drugs. Potentiation occurs with ganglion and peripheral adrenergic blocking agents.

**Sensitivity reactions**  
Sensitivity reactions may occur in patients with a history of allergy or bronchial asthma.

**The possibility of exacerbation or activation of systemic lupus erythematosus has been reported.**

**Use in Pregnancy**  
**Reserpine**  
The safety of reserpine for use during pregnancy or lactation has not been established; therefore, the drug should not be used in pregnant patients or in women of childbearing potential only when, in the judgment of the physician, the potential benefit to the welfare of the patient, increased maternal or neonatal tract secretions, nasal congestion, coryza, and anuria, may occur in neonates and breast-fed infants of reserpine-treated mothers since this drug crosses the placental barrier and appears in breast milk.

**Hydralazine**  
Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. In patients with severe renal disease, the drug may develop in patients with impaired renal function. Thiazides should be used with caution in patients with impaired hepatic function or progressive liver disease, since alterations of fluid and electrolyte balance may be precipitated. Thiazides may be additive or potentiative in the action of other antihypertensive drugs. Potentiation occurs with ganglion and peripheral adrenergic blocking agents.

**Sensitivity reactions**  
Sensitivity reactions may occur in patients with a history of allergy or bronchial asthma.

**The possibility of exacerbation or activation of systemic lupus erythematosus has been reported.**

**Use in Pregnancy**  
**Reserpine**  
The safety of reserpine for use during pregnancy or lactation has not been established; therefore, the drug should not be used in pregnant patients or in women of childbearing potential only when, in the judgment of the physician, the potential benefit to the welfare of the patient, increased maternal or neonatal tract secretions, nasal congestion, coryza, and anuria, may occur in neonates and breast-fed infants of reserpine-treated mothers since this drug crosses the placental barrier and appears in breast milk.

**Hydralazine**  
Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. In patients with severe renal disease, the drug may develop in patients with impaired renal function. Thiazides should be used with caution in patients with impaired hepatic function or progressive liver disease, since alterations of fluid and electrolyte balance may be precipitated. Thiazides may be additive or potentiative in the action of other antihypertensive drugs. Potentiation occurs with ganglion and peripheral adrenergic blocking agents.

**Sensitivity reactions**  
Sensitivity reactions may occur in patients with a history of allergy or bronchial asthma.

**The possibility of exacerbation or activation of systemic lupus erythematosus has been reported.**

**Use in Pregnancy**  
**Reserpine**  
The safety of reserpine for use during pregnancy or lactation has not been established; therefore, the drug should not be used in pregnant patients or in women of childbearing potential only when, in the judgment of the physician, the potential benefit to the welfare of the patient, increased maternal or neonatal tract secretions, nasal congestion, coryza, and anuria, may occur in neonates and breast-fed infants of reserpine-treated mothers since this drug crosses the placental barrier and appears in breast milk.

**Hydralazine**  
Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. In patients with severe renal disease, the drug may develop in patients with impaired renal function. Thiazides should be used with caution in patients with impaired hepatic function or progressive liver disease, since alterations of fluid and electrolyte balance may be precipitated. Thiazides may be additive or potentiative in the action of other antihypertensive drugs. Potentiation occurs with ganglion and peripheral adrenergic blocking agents.

**Sensitivity reactions**  
Sensitivity reactions may occur in patients with a history of allergy or bronchial asthma.

**The possibility of exacerbation or activation of systemic lupus erythematosus has been reported.**

**Use in Pregnancy**  
**Reserpine**  
The safety of reserpine for use during pregnancy or lactation has not been established; therefore, the drug should not be used in pregnant patients or in women of childbearing potential only when, in the judgment of the physician, the potential benefit to the welfare of the patient, increased maternal or neonatal tract secretions, nasal congestion, coryza, and anuria, may occur in neonates and breast-fed infants of reserpine-treated mothers since this drug crosses the placental barrier and appears in breast milk.

**Hydralazine**  
Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. In patients with severe renal disease, the drug may develop in patients with impaired renal function. Thiazides should be used with caution in patients with impaired hepatic function or progressive liver disease, since alterations of fluid and electrolyte balance may be precipitated. Thiazides may be additive or potentiative in the action of other antihypertensive drugs. Potentiation occurs with ganglion and peripheral adrenergic blocking agents.

**Sensitivity reactions**  
Sensitivity reactions may occur in patients with a history of allergy or bronchial asthma.

**The possibility of exacerbation or activation of systemic lupus erythematosus has been reported.**

**Use in Pregnancy**  
**Reserpine**  
The safety of reserpine for use during pregnancy or lactation has not been established; therefore, the drug should not be used in pregnant patients or in women of childbearing potential only when, in the judgment of the physician, the potential benefit to the welfare of the patient, increased maternal or neonatal tract secretions, nasal congestion, coryza, and anuria, may occur in neonates and breast-fed infants of reserpine-treated mothers since this drug crosses the placental barrier and appears in breast milk.

**Hydralazine**  
Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. In patients with severe renal disease, the drug may develop in patients with impaired renal function.



Dr. Lewis and the dialyzer cartridge

## No-pump dialyzer

**T**HE NEW MARKLEY DIALYZER has two important advantages over most others now in use," says Edmund J. Lewis, M.D., director of the University of Chicago's Hemodialysis Unit, where the new artificial kidney is being tested clinically.

"The first is its compactness. At any given time, there are only about 25 cc of blood in the dialyzer—and therefore outside the body—and I think this benefits the patient, particularly the one who is on home dialysis.

"The second advantage, a consequence of the first, is its greater safety over other dialyzers. A majority of our patients will be able to use this unit without a pump, the heart serving as the pump, which simplifies the procedure considerably. The great danger in home dialysis has always been the possibility of a severe blood loss in the event of accident.

"During dialysis, between 300 and 400 cc of blood a minute are pumped out of a patient, and if warning mechanisms remain imperfect, as they are now, the pump may continue to work even when there is a disconnected line, or a ruptured membrane that no one notices.

"But even aside from that hazard, there is a certain amount of trauma to the blood as it goes through the pump. I think patients who are able to use the dialyzer without the pump somehow feel better when the blood flows at its own rate."

Physicist Finley Markley, who developed the new dialyzer at the Argonne National Laboratory in collaboration with University of Chicago physicians, contributed the first departure in dialyzer design when he learned how to bond Cuprophran, the very fine cellophane used

as a membrane in most dialyzers, so that it would be impervious to water.

"This dialyzer has 60 pleats on one side and 61 on the other," said Dr. Lewis. "The blood flows along one side of the pleats while the dialysate flows along the other. The thinner the film of blood you have, the more efficiently molecules of urea and creatinine can move through the cellophane, because they're always in very close contact with the dialysate on either side. That is the key to why this dialyzer is 30 per cent more effective than any others now in use."

Whether a patient will be able to use the Markley unit without a pump depends on the size of his blood vessels and the kind of access available to the vessels. Until recently, the university's Hemodialysis Unit had been using two standard types of arteriovenous shunts. The first, the Scribner shunt—a plastic shunt that comes out of the skin in connection with

**Patients who are able to use the dialyzer without the pump somehow feel better when the blood flows at its own rate.**

the vessels, forming an A-V cannula—is losing popularity because of associated infections, and because of its restraints on patients. They can't swim, and they must be very careful when they bathe.

However, the Scribner shunt can be used with the Markley dialyzer without a pump, because access to the vessels is simply a matter of disconnecting shunt leading between the artery and the vein and plugging into the machine.

The second type of shunt is the so-called internal arteriovenous fistula. To create it, a surgeon connects an artery and a vein in the arm subcutaneously, producing direct communication of blood. The veins become much larger because the artery is feeding them directly rather than going through the capillary bed. Access is made by inserting a needle into a vein in each arm, the shunted one furnishing the arterial supply, the other supplying venous return.

### Infection infrequent

"This is the most convenient kind of access for most patients. It involves no foreign body, no tube coming out of the skin. After a while, they don't mind the insertion of the needles for each dialysis. Many of them do this themselves. But the blood flow we get for the arterial supply is usually not high enough in volume to run the dialysis without a pump."

The unit is now using another approach, the Thomas shunt, which Dr. Lewis believes will make it possible to use the new dialyzer without a pump. The Thomas shunt involves surgical implantation of dacron grafts into a major artery and vein, either in the axillary or femoral area. Each graft is then extended down, to emerge through the skin of the arm or the leg. The larger vessels provide a higher volume of blood supply.

Infection, while still a possible source of trouble in this external access, seems to occur infrequently, probably because connective tissue tends to interpenetrate and seal off the dacron webbing that covers the fistula.

"We have about 35 patients on dialysis now, most of whom are between 20 and 35 years of age and have chronic renal

failure. The typical patient has a history of untreated, or only intermittently treated, hypertension, and developed severe symptoms suddenly, followed by rapid deterioration of kidney function to the point of renal failure and the need for dialysis."

Drop-outs from care in the hypertension clinic are a problem. Some patients just stop taking medication when their blood pressure drops. Even some with renal failure and the ultimate motivation who understand fully the nature of their illness, can find it difficult to maintain the regimen.

Dr. Lewis feels that these patients could have avoided catastrophic illness if their hypertension had been controlled consistently from the start. "I don't mean to indict the patients. There is nothing in the histories we've been able to elicit that contradicts the possibility that physicians failed to prescribe, or that they didn't direct patients to stop medication once blood pressure had been brought down to normal. The situation simply highlights the enormous need for nationwide patient and doctor education.

"If a patient is doing well on dialysis—psychologically, vocationally, and physically—he has a better than 90 per cent yearly chance for survival. The majority of our patients on dialysis are being offered at least 10 more years of life. Who knows what will develop in that time?

"Most of our patients will be kept on dialysis for years before being offered kidney transplants, for they do not have related donors, and results with cadaver transplants, even in the best of hands, are not nearly so good. When it works well

there is no doubt that the patient is better off than he would be on dialysis. But if it doesn't work well, very serious, irreversible problems may occur, even if the kidney is removed and the patient is taken off immunosuppressive drugs."

Right now, a number of scheduled transplant operations are being delayed at the University of Chicago hospitals because there is considerable uncertainty

### The greater danger in home dialysis has always been the possibility of a severe accidental blood loss.

about how the provision of the new Federal law will be interpreted. The law says that a patient must be on a program for three months before the Social Security Department will take over payments. What is unclear is whether patients already on the program now will still have to go through a waiting period, "which only increases morbidity and mortality."

"The LifeMed dialysis delivery machine we use for home dialysis costs somewhere between \$5,000 and \$6,000. Most dialyzer cartridges cost about \$15, and we think the Markley dialyzer will run the same when it goes into commercial production. Salts for the dialysate cost about \$30. Simply for supplies, dialysis costs \$135 a week. And when it is done in a hospital, you must add on the cost of staff and overhead.

"We don't know what is going to happen to these programs. Illinois was once the most advanced state in this area. Five

years ago, before any other state, it enacted legislation to support patients on dialysis. Twenty-eight states have now patterned their programs after that legislation. Criteria for eligibility, however, excluded malignant hypertension, probably because a 30-year-old with chronic renal disease who has had hypertension for several years is at higher risk than another 30-year-old with chronic renal disease and no history of hypertension. Only in the last two years has the state begun to make exceptions.

"This is precisely what we are afraid of in the Federal bill. States had already cut back budgets in expectation of Federal money. But if mechanisms of the Federal law limit payments for dialysis, or set criteria that exclude malignant hypertension, it is going to be a catastrophe, particularly for patients in low socioeconomic groups.

"This kind of legislation can be fiscally prudent but medically imprudent. I don't see a patient as a factor in the GNP or the Federal budget. Yet I may have to face making life or death decisions based upon whether or not a patient qualifies for State or Federal support, or has third-party insurance.

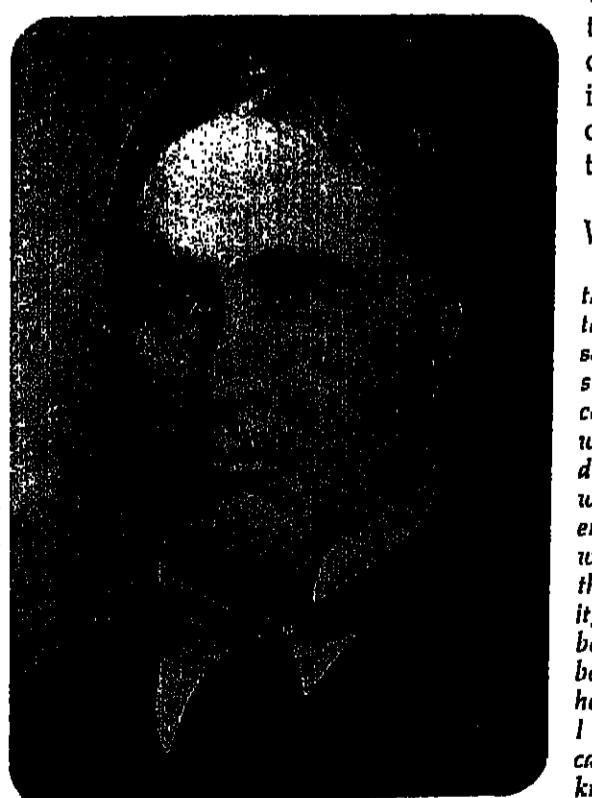
"All the hard work that has gone into getting people to recognize the problem, and getting legislation to cope with it, can be wiped out if government agencies do not take a realistic approach to the payment of expenses of care. Research in transplant surgery of all kinds could slow down and we would lose the gains we've made in recent years. I think it would take the country a long time to recover them, if it ever does."

## HYPERTENSION CLASSICS ... Sir William Gull: the arterioles

**S**IR WILLIAM WITHEY GULL (1816-90) of Guy's, pathologist, clinician and sarcastic epigrammatist ("Make haste and use all medicines before they lose their effectiveness."), presented a paper with H. G. Hutton of London Hospital in 1872, in which they described three forms of chronic Bright's disease.

They found (1) kidneys often much contracted, heart much hypertrophied, minute arteries and capillaries throughout the body thickened by hyalin-fibroid formation; (2) kidneys little contracted, but heart much hypertrophied, minute arteries and capillaries much thickened; (3) kidneys healthy, heart much hypertrophied and minute arteries and capillaries much thickened.

They confirmed George Johnson's findings of alterations in the arterioles (1868), but did not accept that such pathologic change could be attributed to urinary excreta. Rather, it was to be



attributed to a hyalin-fibroid formation in the walls of the minute arteries and a hyalin-granular change in the corresponding capillaries—occurring chiefly outside the muscular layer, but also in the tunica intima of some arterioles throughout the body. "This morbid change in the arterioles and capillaries is the primary and essential condition called chronic Bright's disease with contracted kidney."

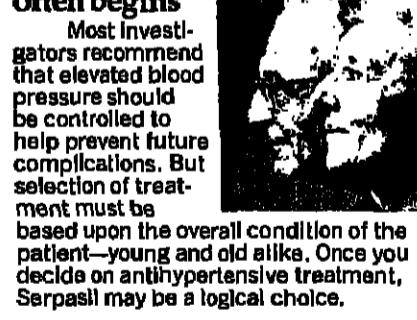
In a lecture given in the same year, Sir William said:

*It is always dangerous to rest in a narrow pathology; and I believe that to be a narrow pathology which is satisfied with what you now see before me on this table. In this glass you see a much hypertrophied heart and a very contracted kidney. This specimen is classical. It was, I believe, put up under Dr. Bright's own direction, and with a view of showing that the wasting of the kidney is the cause of the thickening of the heart, I cannot but look upon it with veneration, but not with conviction. I think, with all deference to so great an authority, that the systemic capillaries, and, had it been possible, the entire man, should have been included in this vase, together with the heart and kidneys; then we should have had, I believe, a truer view of the causation of the cardiac hypertrophy and of the disease of the kidney.*

# The root of antihypertensive therapy



**Serpasil...where antihypertensive therapy often begins**



**Serpasil reduces the "tension" in hypertension**

Serpasil eases the "tension" that plays an important part in many cases of hypertension.

**Warning:** Mental depression, occasionally severe, can occur with use of Serpasil. Discontinue drug at the first sign of depression.

**Serpasil...the antihypertensive to build on**

If you decide to use Serpasil in combination with other antihypertensive agents, lower dosage of these drugs is permitted, minimizing the incidence and severity of their side effects...an important consideration, particularly in treating the older patient.

**Serpasil...a quality reserpine, assured by quality control**

Serpasil, the original reserpine, is established as a quality reserpine. Exact quality control procedures, including 99 tests performed during the manufacturing process, help guarantee its purity, uniformity, and potency.

**Serpasil lowers blood pressure and slows rapid heart rate**

Serpasil acts both on the autonomic and central nervous systems, lowering arterial blood pressure and slowing rapid heart rate.

## Serpasil® (reserpine)

**INDICATIONS**

Based on a review of this drug by the National Institute of Sciences-National Research Council and/or other information, FDA has classified the indications as follows:

### **Effective**

**Oral Serpasil:** Mild essential hypertension; adjunctive therapy with other antihypertensive agents in the treatment of symptoms of hypertension; relief of symptoms in agitated psychotic states (e.g., schizophrenia), primarily in those individuals unable to tolerate phenothiazine derivatives or those who had required antihypertensive medication.

**Parenteral Serpasil (Intramuscular):** Hypertensive emergencies, such as acute hypertension encephalopathy, in which it is necessary to reduce blood pressure rapidly; particularly in those patients unable to accept oral medication, or to control extensive agitation.

**"Possibly" Effective**

**Oral Serpasil:** As an antianxiety agent in chronic anxiety, manic-depressive states, and in management of anxiety and tension associated with neurodermatitis and other dermatoses; as a tranquilizer for psychotherapy in paranoid and manic states or as adjunctive therapy for the treatment of chronic schizophrenia; for tension headaches; and hypotension; for tension headache; for menopausal symptoms; for general paresis; for hypertension of toxemia; for tachycardia and palpitations; for conditions in which there is a history of common hypertension, such as anxiety, tension, nervousness; and for angina pectoris.

**Parenteral Serpasil (Intramuscular):** Psychiatric conditions such as paranoid, manic states, and the manic phase of manic-depressive psychosis; tachycardia; anxiety and tension.

Final classification of the less-than-effective indications requires further investigation.

### **CONTRAINDICATIONS**

Known hypertension, mental depression, or any condition which may induce peptic ulcer, ulcerative colitis, digitalis intoxication, aortic insufficiency, and patients receiving electroconvulsive therapy.

**WARNING:** Mental depression, which may be severe enough to result in suicide, can occur in association with the use of this drug, whether or not there is a previous history of depression or any other psychiatric disorder. If depression occurs, discontinue the drug if the first evidence of depression, such as early morning insomnia, loss of appetite, impotence, or self-deprecating. Extreme caution should be exercised in treating those patients with a history of depression. Drug-induced depression may persist for several months after drug withdrawal.

The drug should be discontinued for at least two weeks before giving electroshock therapy. MAO inhibitors should be avoided or used with extreme caution.

### **Usage in Pregnancy**

The safety of reserpine for use during pregnancy or lactation has not been established; therefore, the drug should be used in pregnant patients or nursing mothers only when, in the judgment of the physician, it is essential to the welfare of the patient.

Increased respiratory tract secretions, nasal congestion, cyanosis, and anoxia may occur in infants. These effects are seen in treated mothers since this drug is known to cross a placental barrier and to appear in breast milk.

**PRECAUTIONS**

Since Serpasil increases gastrointestinal motility and may cause diarrhea, caution is indicated in patients with a history of peptic ulcer, ulcerative colitis, or other gastrointestinal disorders. It may precipitate biliary colic in patients with gallstones.

Because of the effect of catecholamine depletion, amines are more apt to be hyperactive to the drug and their condition may be aggravated. Therefore, special care should be exercised when treating patients with a history of hypertension.

Cautions should be exercised when treating hypertensive patients with renal insufficiency since they adjust poorly to lowered blood pressure levels.

Use with caution with digitalis and quinidine since cardiac arrhythmias have occurred with rawwill preparations.

Concurrent use of guanethidine and rawwill derivatives may cause bradycardia, mental depression, and postural hypotension. Hypertension may persist at a higher risk of intrapertoneal hypertension and other cardiovascular complications than normotensive patients. Serpasil-treated patients are not known to have a higher risk of such complications than otherwise comparable hypertensive patients.

Preoperative withdrawal of reserpine does not assure that circulatory instability will not occur. It is important that the anesthesiologist be aware of the patient's drug intake and consider this in the overall management, since hypertension has occurred in patients receiving reserpine, guanethidine, and/or adrenergic and/or adrenergic drugs (e.g., mescaline, morphine, neophrine) have been employed to treat adverse vagocirculatory effects.

### **ADVERSE REACTIONS**

Adverse reactions have caused gastrointestinal reactions including hyperacidity, nausea, vomiting, anorexia, and diarrhea; cardiovascular reactions particularly when used concomitantly with digitalis or quinidine; and bradycardia, central nervous system reactions including drowsiness, depression, nervousness, paradoxical anxiety, nightmares, and, rarely, parkinsonian syndrome and other extra-pyramidal side effects; and CNS sensitization manifested by hallucinations, delusions, glaucoma, uveitis, and optic atrophy. Nasal congestion is a frequent complaint. Pruritus, rash, dryness of mouth, dizziness, headache, dyspnea, syncope, epistaxis, purpura and other hematological reactions; and, in addition to drowsiness, dysuria, muscular aches, conjunctival injection, weight gain, breast engorgement, pseudogout, and gynecomastia have been reported.

These reactions are usually reversible and disappear when the drug is discontinued. Nasal congestion is a frequent complaint. Pruritus, rash, dryness of mouth, dizziness, headache, dyspnea, syncope, epistaxis, purpura and other hematological reactions; and, in addition to drowsiness, dysuria, muscular aches, conjunctival injection, weight gain, breast engorgement, pseudogout, and gynecomastia have been reported.

### **DOSAGE & ADMINISTRATION**

#### **Oral Serpasil**

**For Hypertension:** In the average patient not receiving other antihypertensive agents, the usual initial dose is 0.5 mg daily for 1 or 2 weeks. For patients with hypertension, 0.25 mg daily. Higher doses should be used cautiously, because serious mental depression and other side effects may be increased considerably.

Serpasil may be combined with other antihypertensive agents—such as a thiazide and/or chlorothiazide, to bring about a maximal therapeutic response.

**For Psychiatric Disorders:** The usual initial dose is 0.5 mg orally with a range of 0.1 to 1.0 mg. Adjust dosage upward or downward according to the patient's response.

**For Tachycardia:** Recommended dosage range is 0.1 to 0.5 mg orally per day. Rapid heart rate is to be expected, up to 2 times the normal rate at which time the drug dose should be reduced.

Suppression of tachycardia often persists after therapy is stopped. Note: In patients receiving digitalis or quinidine, give reserpine cautiously. It is not recommended in cases of digitalis or quinidine.

**For Anxiety-Tension and Related Disorders:** Initial daily dosage range is 0.1 to 0.5 mg orally, as a single dose or in divided doses. For maintenance, adjust dosage according to response. For children, as little as 0.1 mg per day is often sufficient.

**Parenteral Serpasil (Intramuscular):** Serpasil may be administered parenterally in the short-term treatment of hypertensive crisis. Initial or emergency oral or parenteral reserpine should be used. An initial dose of 0.1 to 1 mg intramuscularly is followed at intervals of 3 hours, if necessary, by doses of 2 and 4 mg until the blood pressure falls to the desired level. If the 4-mg dose is ineffective, other antihypertensive agents should be used. An initial dose larger than 0.5 mg may induce severe hypotension, particularly in patients with cerebral hemorrhage.

Cardio titration of dosage is required for combination therapy with Serpasil with other antihypertensive agents.

Serpasil may be administered intramuscularly in psychiatric emergencies to initiate treatment in those patients unable to accept oral medication or to accept oral agents. The initial dose is from 2.5 mg to 5.0 mg, following a small initial dose to test sensitivity.

**HOW SUPPLIED**

Tablets, 1 mg (white, scored); bottles of 100, 500, 1,000, 5,000 and Strip Dispensers of 100.

Tablets, 0.1 mg (white); bottles of 100, 500 and 1,000.

**Elixir** (green, lemon-lime flavored), 0.2 mg per 5 ml; bottles of 100.

**Parenteral:** Each ml contains 2.5 mg reserpine, 0.1 ml dimethylacetamide, 10 mg adipic acid, 0.1 mg vorscine, 0.01 ml benzyl alcohol, 0.05 ml polyethylene glycol, 0.5 mg ascorbic acid, and 0.1 mg sodium sulfite. In 1 ml Ampoules, 2 ml cartridges, 500. Multiple-dose Vials, 10 ml; cartons of 1, boxes of 6.

Rev. 3/72

**CIBA Pharmaceutical Company**  
Division of CIBA-GEIGY Corporation  
Summit, New Jersey 07901

# Serpasil (reserpine)

early effective control of hypertension can save lives

CIBA

C/3411M

Wednesday, August 8, 1973

MEDICAL TRIBUNE

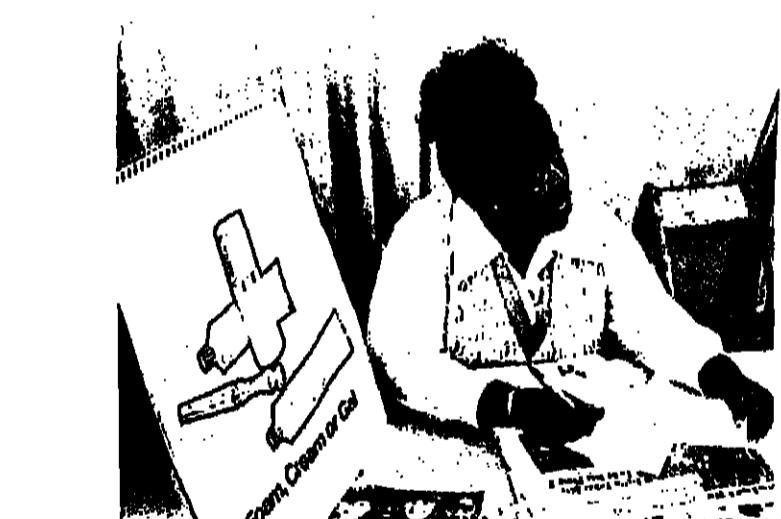


Medical Tribune Photo—Harry Rhine

Above at left, elderly persons enjoy their ride up the Hudson. The program run by the Hospital Ship gave the participants a day's outing and lunch as well as several types of screening, care, and advice. At right, one of the women has her blood pressure taken. Eye examinations, blood tests, and general physical examinations were among the tests also given. The Hospital Ship program reaches several thousand of the area's senior citizens each summer.



Many elderly have no access to a doctor and no idea where to go for care.



At the Brookdale Fair, held in a striped tent, the emphasis was not only on examinations but also on education and advice. Many of the street health fairs, like the one at right, are held with the assistance of local block groups.



At the Brookdale Fair, refreshments, bands, a puppet show, and special attractions such as the trampoline helped in relaxing the children.

Street fairs provided measles and rubella immunizations and tests for lead poisoning, sickle cell anemia, and VD. Hypertension screening, drug referrals, and lectures on dental care were also given.



## Clinical Trials



## Higher Mortality Seen in Teen Pregnancies

Medical Tribune Report

ANN ARBOR, MICH.—The death rate for teen-age mothers and their babies is 30 per cent higher than that for mothers age 20 to 24 years and their babies, a Detroit physician reported here, asserting that the situation "is not just fetal and maternal wastage."

"There are large social consequences in teen-age pregnancy—curtailed education, the mire of economic disadvantage, the emotional conflict, and perhaps suicide," Dr. Joan Stryker, of Hutzel Hospital, told a seminar on human sexuality at the University of Michigan.

Dr. Stryker, who is also medical director of the Planned Parenthood League of Detroit, said that there is a great need to provide teen-agers with more intensive, accurate, and honest information on all aspects of sex.

She described the early results of two birth control centers for teen-agers that she helped to establish. One is a clinic for pregnant girls 17 years old and younger, operated at the hospital, and the other a Planned Parenthood teen center for both boys and girls.

Describing "subtle" changes in the two years that the clinics have been operating, she said that, first, there are fewer irate parents calling, "in fact, the parents are becoming involved even to assistance in the form of money and time." Another change, she reported, has been in education—the patients request more sex information and the word has gone out that "the clinics offer worthwhile information and comprehensive health services."

A survey of the pregnant girls in the clinics, conducted this spring, found that 81 per cent had never used a contraceptive; in a survey conducted last fall the figure was 95 per cent.

## 64 % Requested Abortions

Both surveys showed that about 64 per cent of the girls requested abortions. Of the 36 per cent who chose to deliver their babies, the spring survey disclosed that about half said they intended to raise their babies and remain single, 15 per cent said they intended to marry, and 3 per cent said they would place the infants for adoption. In the fall survey, the respective figures were 5 per cent, 21 per cent, and 9 per cent.

Describing her approach to sex counseling and the provision of contraceptives, Dr. Stryker said, "If the teen-ager has never been sexually active but comes for contraceptive advice, I personally talk to

her for two reasons—the legal, so that we cannot be accused of contributing to the delinquency of a minor, and to be sure the patient really wants to have sexual intercourse. Subconsciously, they may want a brake. Some girls are relieved when I say No." She noted that "the door is left open a crack" and the patient is assured that when she is ready to have intercourse she could come back.

With respect to the contraceptives prescribed, Dr. Stryker said that for the girl who has never been pregnant and who has intercourse less than once a month, it is suggested that she use vaginal jelly and her partner a condom. For the more sexually active girls, birth control pills are prescribed for those in good hormonal balance, but a diaphragm, intrauterine device, or combination of jelly and condom for those with irregular menstrual cycles.

Dr. Stryker recommended the pill for girls who have been pregnant, but she added that an intrauterine device may be a better choice of contraceptive for young women who are likely to forget to take the pill regularly.

ally active girls, birth control pills are prescribed for those in good hormonal balance, but a diaphragm, intrauterine device, or combination of jelly and condom for those with irregular menstrual cycles.

Dr. Stryker recommended the pill for girls who have been pregnant, but she added that an intrauterine device may be a better choice of contraceptive for young women who are likely to forget to take the pill regularly.

# Gastroenterology

## Proper Training, Temperament Urged in Use of Colonoscope

Medical Tribune Report

New York—A warning that anyone planning to use the new fiberoptic colonoscopes should have adequate training as well as the right kind of temperament was sounded during a panel discussion of colonoscopy here at the annual convention of the American Medical Association.

Use of these highly sophisticated instruments is now increasing rapidly, yet almost no courses are available to provide instruction in techniques, the panel members told a session held jointly by the Section on Colon and Rectal Surgery and the Section on Gastroenterology.

"Fiberoptic colonoscopy constitutes a real advance in the diagnostic and therapeutic armamentarium of the medical profession," said Dr. William I. Wolff, of New York's Beth Israel Medical Center, where he and colleagues have performed more than 4,000 diagnostic colonoscopic examinations and some 600 polypectomies.

The colonoscope amounts to a probe, he said, quoting the old surgical maxim that "there are few more dangerous instruments than a probe with no brains behind it."

By Dr. Wolff's definition, competence includes the following:

• Manual dexterity, termed essential.

- An interest in what is being done.
- Knowledge of gastrointestinal diseases, since what is seen "must be correctly interpreted and put into context with the patient's clinical symptomatology."
- Suitable temperament. It takes time "to develop the skill required to pass the instruments and to do this safely—if the physician is going to hurry, he's bound to get into trouble."

• Judgment. "The endoscopist must recognize his limitations, the limitations of the case, and when to proceed and when not to go ahead."

A fellow panelist and surgeon, Dr. Howard Jay Eddy, Jr., of Garden City, Long Island, also advocated setting up workshops or courses to provide training.

"With the legal implications of performing colon surgery without a prior colonoscopy looming on the horizon, the necessity for the establishment of training programs is of the greatest importance," he said.

Meanwhile, Dr. Eddy thinks some form of limitation should be imposed on the "would-be instant endoscopist."

Discussing reported instances in which physicians have bypassed offers of training because they preferred to "try out procedures for themselves," he said he had only one comment:

"God help the patient and the procedure!"

# Duodenal Ills May Be Causing Upper GI Bleeding

Medical Tribune Report

New York—Hemorrhagic duodenitis, an entity distinct from peptic ulcer disease or other known inflammatory diseases of the esophagus or stomach, should be considered in the differential diagnosis of upper gastrointestinal bleeding, a Michigan gastroenterologist emphasized here.

Of 35 patients with nonspecific duodenitis, eight were observed with hemorrhagic duodenitis significant enough to cause acute upper gastrointestinal bleeding as evidenced by hematemesis and/or melena and blood-loss-type anemia, Dr. Eugene A. Gelzayd of Southfield, Mich., told a meeting of the American Society for Gastrointestinal Endoscopy, held in conjunction with the annual meeting of the American Gastroenterological Association.

Clinically, he reported, anemia was present in seven of the eight patients, and four required multiple transfusions. Hematemesis was present in four cases and melena in all eight. Dyspeptic epigastric distress in five patients preceded the onset of the bleeding by a variable period of time. Two patients were asymptomatic and a third had onset of massive bleeding during a myocardial infarction.

Radiologic findings included spasm and irritability of the bulb and coarse folds in the proximal duodenum in six of seven patients studied, said Dr. Gelzayd. Maximum histologic gastric analyses, quantitative serum immunoglobulins, and direct examination of duodenal secretions for Giardia and other parasites were normal in five patients.

## Duodenoscopy Was Abnormal

Duodenoscopy was abnormal in all eight patients and included hyperemia, mucosal friability, and superficial erosions with active bleeding. In three patients there was also mild mucosal nodularity of the bulb and proximal duodenum. Endoscopically, the stomach was normal in all patients. Mild erosive esophagitis without visible bleeding was seen in one acute alcoholic patient; two others with alcoholic liver disease had intact esophageal varices.

Histologic evidence of duodenitis was found in six patients who underwent target mucosal biopsies of the bulb and proximal duodenum, Dr. Gelzayd said. This included superficial ulcerations of the epithelial lining, mild degrees of villous flattening, crypt dilatation, and dense acute and chronic inflammatory cell infiltration of the lamina propria, which, in two patients, extended into the Brunner's gland area. Edema, dilated and engorged capillaries, and intramural hemorrhage were also noted.

A history of significant alcohol intake was recorded in four patients, and aspirin usage before bleeding, averaging four to six tablets a day for several days, in two patients.

Conservative treatment, consisting of nasogastric intubation and suction, antacids and milk, diazepam, and reassurance, said Dr. Gelzayd, resulted in cessation of bleeding in seven patients for up to two years' follow-up. The patient with the myocardial infarction died.

"Correction of anemia with blood transfusion or iron is also important," he said.

He added that, "conceivably, surgery may be advisable in some cases for uncontrollable hemorrhage."

He recommended endoscopic follow-up in patients with hemorrhagic duodenitis. Follow-up duodenoscopy and biopsy were performed in five of the patients, nine to 18 months after the initial bleeding episode. Three demonstrated mild patchy hyperemia and friability, one mild nodularity of the bulb mucosa, and the fifth a normal-appearing duodenum. Actively bleeding erosions were not seen, but in all five patients histologic, inflammatory changes of duodenitis were still present.

Coauthor of the report was Dr. David Gelfand.

# the bare facts

in many dermatoses the less they wear, the more they need...

## Vioform-Hydrocortisone (dodechlorhydroxyquin and hydrocortisone)

antifungal • antibacterial • anti-inflammatory • antipruritic

Some styles don't leave much to the imagination. And don't provide much cover for common dermatoses, either. Just like plain topical steroids. If the lesion has become infected with fungi or bacteria, plain topical steroids are ordinarily not recommended as sole therapy. Vioform-Hydrocortisone, on the other hand, provides the kind of comprehensive therapy these dermatoses may require. It not only supplies the anti-inflammatory and antipruritic actions of hydrocortisone... but also adds the antibacterial and antifungal actions of Vioform.

\*This drug has been evaluated as possibly effective for these indications. See brief prescribing information.

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream

Another fact...  
the most widely  
prescribed form...  
20 Gm cream



If there's good reason  
to prescribe  
for psychic tension...



When, for example, despite counseling, tension and anxiety continue to produce distressing somatic symptoms

Prompt action  
is a good reason  
to consider Valium®  
(diazepam)

When your patient's somatic complaints are associated with tension and anxiety and you have tried counseling and other supportive measures alone, you may decide to prescribe psychotherapeutic medication. If you do, the question remains: which one?

Valium (diazepam) is one to consider closely. One that works promptly as an adjunct to continued supportive measures. One that generally produces significant improvement within the first few days of therapy, although some patients may require more time for a clear-cut response.

Prompt action. One good reason to consider Valium.

And should you choose to prescribe Valium, you should also keep this information in mind. Valium is usually well tolerated; the most common side effects reported have been drowsiness, fatigue and ataxia. Patients taking Valium should be cautioned against operating dangerous machinery or driving. Therapy with Valium should normally be continued until the patient's psychic tension symptoms have been reduced to tolerable levels.

Please turn page for a summary of product information.

**Valium®**  
(diazepam)

2-mg, 5-mg, 10-mg tablets

ROCHE

# Other good reasons to consider Valium® (diazepam)

## Effectiveness

The efficacy of Valium (diazepam) has been proven in clinical studies and in extensive clinical use. It can relieve psychic tension and its somatic symptoms in patients who overreact to stress and in psychoneurotic patients.

**Before prescribing, please consult complete product information, a summary of which follows:**

**Indications:** Tension and anxiety states, somatic complaints which are concomitants of emotional factors; psychoneurotic states manifested by tension, anxiety, apprehension, fatigue, depressive symptoms or agitation; symptomatic relief of acute agitation, tremor, delirium tremens and hallucinosis due to acute alcohol withdrawal; adjunctively in skeletal muscle spasm due to reflex spasm to local pathology, spasticity caused by upper motor neuron disorders, athetosis, stiff-man syndrome, convulsive disorders (not for sole therapy).

**Contraindicated:** Known hypersensitivity to the drug. Children under 6 months of age. Acute narrow angle glaucoma; may be used in patients with open angle glaucoma who are receiving appropriate therapy.

**Warnings:** Not of value in psychotic patients. Caution against hazardous occupations requiring complete mental alertness. When used adjunctively in convulsive disorders, possibility of increase in frequency and/or severity of grand mal seizures may require increased dosage of standard anticonvulsant medication; abrupt withdrawal may be associated with temporary increase in frequency and/or

## Dependable response

The psychotherapeutic effect of Valium (diazepam), characterized by symptomatic relief of tension and anxiety, is generally reliable and predictable.

## Titratable dosage

With Valium (diazepam), adjustments in dosage can alter the clinical response. This titratability enables you to tailor your therapy for maximum efficiency. There are three convenient tablet strengths to choose from: 2 mg, 5 mg and 10 mg.

severity of seizures. Advise against simultaneous ingestion of alcohol and other CNS depressants. Withdrawal symptoms (similar to those with barbiturates and alcohol) have occurred following abrupt discontinuance (convulsions, tremor, abdominal and muscle cramps, vomiting and sweating). Keep addiction-prone individuals under careful surveillance because of their predisposition to habituation and dependence. In pregnancy, lactation or women of childbearing age, weigh potential benefit against possible hazard.

**Precautions:** If combined with other psychotropics or anticonvulsants, consider carefully pharmacology of agents employed; drugs such as phenothiazines, narcotics, barbiturates, MAO inhibitors and other antidepressants may potentiate its action. Usual precautions indicated in patients severely depressed, or with latent depression, or with suicidal tendencies. Observe usual precautions in impaired renal or hepatic function. Limit dosage to smallest effective amount in elderly and debilitated to preclude ataxia or oversedation.

**Side Effects:** Drowsiness, confusion, diplopia, hypotension, changes in libido, nausea, fatigue, depression, dysarthria, jaundice, skin rash, ataxia, constipation, headache, incontinence, changes in

salivation, slurred speech, tremor, vertigo, urinary retention, blurred vision. Paradoxical reactions such as acute hyperexcited states, anxiety, hallucinations, increased muscle spasticity, insomnia, rage, sleep disturbances, stimulation have been reported; should these occur, discontinue drug. Isolated reports of neutropenia, jaundice; periodic blood counts and liver function tests advisable during long-term therapy.

**Dosage:** Individualize for maximum beneficial effect. **Adults:** Tension, anxiety and psychoneurotic states, 2 to 10 mg b.i.d. to q.i.d.; alcoholism, 10 mg t.i.d. or q.i.d. in first 24 hours, then 5 mg t.i.d. or q.i.d. as needed; adjunctively in skeletal muscle spasm, 2 to 10 mg t.i.d. or q.i.d.; adjunctively in convulsive disorders, 2 to 10 mg b.i.d. to q.i.d. **Geriatric or debilitated patients:** 2 to 2½ mg, 1 or 2 times daily initially, increasing as needed and tolerated. (See Precautions.) **Children:** 1 to 2½ mg t.i.d. or q.i.d. initially, increasing as needed and tolerated (not for use under 6 months).

**Supplied:** Valium® (diazepam) Tablets, 2 mg, 5 mg and 10 mg; bottles of 100 and 500. All strengths also available in Tel-E-Dose® packages of 1000.

**ROCHE**  
Roche Laboratories  
Division of Hoffmann-La Roche Inc.  
Nutley, N.J. 07110

## Physicians Are Blamed for 'People Pollution'

*Medical Tribune World Service*  
AUCKLAND, NEW ZEALAND—A scientist and a former cabinet minister in the New Zealand Government have blamed doctors for "people pollution."

Speaking at the Medical Association of New Zealand, former Minister for the Environment Duncan MacIntyre told the doctors: "Whether you like it or not, one of the problems of environmental control which will be dumped in your lap is that of euthanasia. It is your duty to advise us in how to control the number of people."

Mr. MacIntyre said the world population explosion, which young people are calling the "ultimate pollution," presents many problems for the medical profession.

He asserted that physicians are responsible for much of it, with the millions of lives saved by drugs and the extension of men's lives by money and skill.

The scientist, Ian L. Baumgart, who is assistant director-general of the Department of Scientific and Industrial Research

in New Zealand, told the conference: "The real cause of the problems we are facing is not, fundamentally, advancing technology or rising standards of living or willful destruction of natural resources."

"The fundamental cause is people, more and more of them brought into the world by the medical profession, kept alive to breed by the medical profession, kept alive

to age . . . and to make a bigger impact on all stages of life by the medical profession."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

Mr. Baumgart said scientists were sick

of being blamed for creating problems

that were caused primarily by the ever-

increasing world population, "for which

you medical people are primarily respon-

sible."

&lt;p



### Hospital Provides No Escape From School

In a school on the pediatrics wing of West Virginia University Hospital, volunteer tutor Ann Morgan assists patients studying arithmetic. An elementary school teacher for four years, Mary Ellen Smith (at board) conducts classes for patients ranging in age from prekindergarten to junior high.

## British Psychiatrist Urges Lithium Use for Depression

Medical Tribune World Service

ST. MORITZ, SWITZERLAND—The treatment of depression has been transformed by the introduction of a range of successful drugs, speakers at an international symposium agreed here. In an interview with MEDICAL TRIBUNE a British psychiatrist—Dr. Alec Coppen, of the Medical Research Council—made a plea for an even wider approach to depression, based on prophylaxis with lithium. Work on these lines in the United Kingdom has demonstrated that it can bring about a marked fall in morbidity, Dr. Coppen declared.

MEDICAL TRIBUNE Dr. Coppen, society is becoming increasingly aware that depression is a health problem on a scale comparable to diabetes or leukemia. How far have we advanced in therapy?

Dr. Coppen: The advent of the tricyclic antidepressants has meant that we can now treat depression successfully. However, my view is that we should be working much further back along the line and seeking to prevent depressive episodes.

M.T.: You refer to lithium therapy?

Dr. Coppen: Yes. If prophylactic methods

based on lithium are applied properly, we could hope to reduce the morbidity of this illness by about 80 per cent.

M.T.: You speak of prophylaxis. Is this not also really therapy?

Dr. Coppen: The answer to that is linked with the natural history of depression, considered as a disease. There is first a period which, if untreated, lasts several months. An interval occurs, and then along comes a second episode, again lasting several months. We know from epidemiological studies by Angst and others that the free interval tends to shorten with each attack. Thus, there might be a five-year interval between the first episode and the second, but between the second and the third the gap may narrow to three years, and so on. By the time the patient has had three or four attacks, we face not only the morbidity itself but also the question of recurrence at intervals which are now of only a few months.

M.T.: So there are two problems?

Dr. Coppen: In a way, it is all one problem. We can get to the point where we have difficulty in deciding whether an episode is new or part of the previous event. I always try to emphasize that we are dealing with a long-standing recurrent condition, and we must look at treatment in this perspective.

M.T.: In terms of results, how do the conventional and preventive treatments compare?

Dr. Coppen: Let me give you an example. We have been following up patients who have had three or more attacks of depression and were being treated by conventional methods in different centers in the United Kingdom. We found that they were spending, on average, nearly 50 per cent of their time with an episode, which is a very unsatisfactory state of affairs. With a prophylactic or stabilizer such as lithium, we could expect to bring this down to about 9 per cent.

M.T.: The reduction you quote is based on your studies?

Dr. Coppen: Yes. We have a lithium clinic in our unit, and similar work is also being done at other units in the U.K. The patients are seen every six weeks, and their clinical state, including blood plasma levels of lithium, is monitored.

M.T.: Do you have many backsliders?

Dr. Coppen: Our answer to this is to get the patients interested in their blood plasma levels, to get them to see it all as a cooperative effort.

M.T.: What is the longest period any of your patients has been on lithium as a stabilizer?

Dr. Coppen: I have been treating one patient for five and a half years. All the data suggest that certain patients are and will be increasingly vulnerable to a recurrence of depression for the rest of their lives. Our studies have shown that when they are taken off lithium and put on placebo they will often relapse.

M.T.: So they have to be maintained on therapy rather like hypertension or diabetes?

Dr. Coppen: That is the position. It is a long-term task, in which the patient himself must become engaged.

10 per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

Follow-up: A check on the car's ability to hold special tune and to insure no driver factor, a random check made after two weeks.

Ten per cent increased efficiency with tune-up, or put another way, 10 per cent savings on gasoline bills. Impressive? Not if the car was "out of tune," but this car was in tune. It was to manufacturers' standards and was "in tune" when adjustments were made.

#### Illegal Adjustments?

What was done might be in violation of certain state emission standards, although no auto pollution devices were altered or removed. Quite simply, adjustment of air-gas mixture with spark advance for maximum efficiency was carried out. This may allow slightly more hydrocarbon loss, but

Test car: Two-year-old, four-door sedan, automatic transmission, full power, plus air conditioning.

Test mileage: Professional travel to hospital, office, calls, and home.

## TRIBUNE SPORTS REPORT

**Ballplayer Gets Out of Slump Thanks to Hypnosis Session***Continued from page 1*

average has jumped from .258 to .321, making him one of the top hitters in the American League. "I went out to see the Orioles play last night," Dr. Conn told MEDICAL TRIBUNE in an exclusive interview. "You should see him hit that ball." Meanwhile the Baltimore *News American* sports editor is calling for Dr. Conn to get the "MVP" award—"most valuable psychiatrist."

**A Hypnosis Fan**

Are you a baseball fan? MEDICAL TRIBUNE asked Dr. Conn. "No, I'm not a fan of baseball. I'm a fan of hypnosis," said Dr. Conn. "Because this is a beautiful demonstration of what hypnosis is all about. Many physicians forget that hypnosis is one of the oldest forms of treatment known to man. Greek medical records show it was being practiced 500 years B.C. in the temple of Aesculapius. It was also practiced in the ancient healing temples of India."

Dr. Conn, a professor emeritus at Johns Hopkins School of Medicine who still lectures there on hypnosis, is one of the most expert practitioners of hypnosis in the nation. He told MEDICAL TRIBUNE, because of its physician readers, how Blair became his patient and explained why hypnosis became effective in Blair's case: Dr. Conn, a past president of the Society for Clinical and Experimental Hypnosis, was awarded a gold medal for his contributions to hypnosis in 1970.

**How Case Arose**

Blair was an exceptionally fine Oklahoma athlete, a former Golden Gloves champion, well coordinated and conditioned. However, three years ago he was struck by a "beam ball" thrown by Ken Tatum, of the California Angels. The hard-thrown ball struck him in the face and fractured his nose. After that, Blair became nervous at bat. If a ball came close to him, he ducked. He worried about whether he could duck in time. Pitchers began to pitch balls close to him, causing

him to "bail out"—drop to the ground. His heart would pound and he would feel himself go slack.

Blair's batting average slumped. For two seasons he tried to pull himself together, but his nervousness—and his slump—continued. He was slowly being driven out of baseball by fear.

At this point a Baltimore *News-American* sports writer, Chan Keith, who had become a friend of Blair's, sought out that paper's medical writer, Joann Rogers. He asked who could treat fear. Months earlier she had written about Dr. Conn's work in hypnosis, and she now recommended him.

Keith suggested to Blair that he see Dr. Conn. At that point, June 15, Blair had scored only 38 hits and one home run, giving him an average of .258—and the season was one-third over. Blair, desperate, said, "I'll try anything."

Dr. Conn described the treatment for MEDICAL TRIBUNE because "hypnosis isn't well understood and in recent months research psychologists have been challenging it, calling it mere role-playing-like giving an actor a role to play and coaching him in it. Actually, the Blair case is a beautiful example of how hypnosis works."

**Role Playing Comes First**

"Role playing is indeed the first stage. We all do this when we go to the theater and are drawn to some character and begin to feel and react with him through our identification."

"The second stage is regression. In this stage the patient moves back to childhood, to a secure and safe relationship in which

"But now, instead of a kid pitching to him, we move up in fantasy to the father of one of these kids he's been playing with—a man who is a bush-league pitcher himself. And Blair feels safe and protected against danger—and he can hit. So we let him pitch—and Blair can remember going through the bush leagues and the fun he had there and hitting the ball out of the park."

"And all that gives him back his confidence, his way of feeling about himself. I tell him you don't have to be afraid. Just keep your eye on the ball. Your body will take care of you. You're ready for anything."

Blair had only one session with Dr. Conn. He returned immediately to play. At Dr. Conn's suggestion, he sat down before each game, looked at a spot on the ball, counted to 20, closed his eyes, and then recalled all the fun he had playing

one to four years, 214 mg./100 ml.; five to nine years, 222 mg./100 ml.; 10-14 years, 227 mg./100 ml.; and 15-20 years, 23 mg./100 ml.

In discussion, Dr. Pearson said that any recommendation to the Ministry of Health on universal screening of children should await further study. "It may be a good thing and it may not be," he said. "We'll have to wait till we know what effect serum cholesterol has on atherosclerosis."

The Toronto study showed a significant rise in mean serum cholesterol from birth to five years of age and then a leveling off. There were no significant differences by sex. The levels were determined for 2,639 children. After exclusion for secondary causes of hyperlipoproteinemia, a total of 1,232 were left in the study.

In those with levels two standard deviations above the mean, the assays were repeated three times in the fasting state, in addition to assays of serum triglycerides and a lipoprotein electrophoresis, before the diagnosis was confirmed.

The study showed the following upper limits of normal levels for various age groups: zero to three months, 175 mg./100 ml.; four to 11 months, 195 mg./100 ml.;

## MEDICAL MEETING SCHEDULE

**Foreign Meetings**

Sept. 8-9 ... International Congress of the International League Against Epilepsy, Barcelona, Spain  
 Sept. 8-15 ... International Congress of Chemotherapy, Athens  
 Sept. 10-13 ... European Conference on Pediatric Nephrology, Strakova Plesa, Czechoslovakia  
 Sept. 10-13 ... International Symposium on Electro-Response Audiometry, Bordeaux, France  
 Sept. 10-14 ... International Atomic Energy Agency Symposium on Radiolabelled Peptides, Clinica Medica and Research, Istanbul, Turkey  
 Sept. 10-15 ... International Congress on Cybernetics, Nijmegen, Belgium  
 Sept. 12-15 ... Dutch Orthopaedic Association International Congress on the Knee, Joint, Rotation, Balkan Medical Union Meeting, Ankara  
 Sept. 16-20 ... European Symposium on the Impact of Ecological Factors on Peripheral Vascular Disease, International Congress of Rheumatology, Tel Aviv, Israel  
 Sept. 27- Oct. 6 ... Conference on the Pathophysiology of Neuromuscular Function, Cremona, Italy  
 Oct. 6-10 ... International Congress of Rheumatology, Kyoto, Japan

**Boxing Is Down for the Count With Physicians in Australia***Medical Tribune World Service*

CANBERRA—The Australian Government plans to conduct a thorough inquiry into the promotion and control of boxing. It will also consider the establishment of a Federal boxing commission.

The Federal Minister for Recreation, Frank Stewart, announced the inquiry after a call by the Australian Medical Association for stricter controls to safeguard the health of boxers.

He said the government inquiry will range over all aspects of the sport, including promotion and telecasting.

An interdepartmental committee will study:

- Controls already existing in amateur and professional boxing.
- The justification of external controls.
- The possible spheres of influence of a boxing commission and its jurisdiction over participants, trainers, referees, and judges and over stadiums and training facilities.
- The provision of proper medical supervision.
- The provision of adequate insurance and welfare services for boxers, where necessary.
- The legal responsibility of promoters and others in contracts with boxers.
- Any special provision for televised contests.

*Medical Tribune World Service*

SYDNEY, AUSTRALIA—The Federal Council of the Australian Medical Association has called for strict controls on amateur and

Wednesday, August 8, 1973

MEDICAL TRIBUNE

**Acupuncture's No Mystery, Pain Expert Tells A.M.A.***Medical Tribune Report*

Four male monkeys were first driven batty by being kept in total isolation for six months after their birth, then were brought to "virtually complete recovery" by being permitted contact with immature females, three months younger than the males, the University of Wisconsin reports.

The scientists who conducted the experiment refer to these female monkeys as "therapists" (their quotes, incidentally), although, as far as we can make out, the ladies have never been admitted to a clinical program or even begun a training analysis. Their chief claim to credentials is that stimulation of a point may affect a distant area, and the fact that acupuncture effects may persist even after needles are withdrawn.

In another comment, an official of the Food and Drug Administration, David Link, observed that there has never been any "medical technique that has been so highly publicized [as acupuncture] and about which so little is known." He added that there are great opportunities for the unscrupulous to abuse the technique.

He noted that the FDA legal has no power to control over acupuncture instruments or any other medical device. The agency, however, does have authority over "adulteration or misbranding" of these devices, he said. Current FDA policy is to regard as misbranded any claim for diagnostic or therapeutic effectiveness made for any acupuncture device. They must be clearly labeled as experimental, Mr. Link said.

Panel moderator, Dr. Walter Judd, a gate control theory of pain, said that a first step in explaining acupuncture is to drop the older, oversimplified view that there is a simple one-to-one relationship be-

tween stimulus and pain. Transmission of pain, he said, is a dynamic process, capable of being modulated.

"Every culture on every continent has developed its own type of acupuncture," said Ronald Melzack, Ph.D., Professor of Psychology at McGill University, Montreal. He cited scarification, trepanation, and cupping as other pain-relieving techniques that may possibly work by ways similar to acupuncture.

There are three features in acupuncture pain treatment and "not one of them is a mystery to use," he commented. These include hyperstimulation analgesia, the fact that stimulation of a point may affect a distant area, and the fact that acupuncture effects may persist even after needles are withdrawn.

At any rate, the release says they were "too young to be aggressive as peers, or to show behavior more complex than clinging and simple playing." It also notes that in earlier experiments, attempts to rehabilitate isolates by exposing them to normal peers had not worked.

In the treatment plan, the isolate and his teeny-bopper therapist interacted as a pair for two hours a day, three times a week—there was also group therapy. The first response of the therapists to the troubled isolates was just to cling to them in a very accepting way. After about a week, the males returned the clinging—and there's no need for all that giggling in the back row.

Now you can't tell the isolates from the therapists, but the patients, says the study, "cannot be considered completely normal until they become old enough to mate."

The moral of the experiment seems to be that no matter what happens to you, "the potential for recovery remains as long as an appropriately designed teaching method is available to tap this potential."

If we get word of a human experiment along similar lines, we'll report it.

*Medical Tribune Report**Medical Tribune Report*